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(54) Title: A METHOD FOR THE PROPHYLAXIS AND/OR TREATMENT OF MEDICAL DISORDERS

(57) Abstract: The present invention relates generally to a method for the prophylaxis and/or treatment of skin disorders, and in particular proliferative and/or inflammatory skin disorders, and to genetic molecules useful for same. The present invention is particularly directed to genetic molecules capable of modulating growth factor interaction with its receptor on epidermal keratinocytes to inhibit, reduce or otherwise decrease stimulation of this layer of cells. The present invention contemplates, in a most preferred embodiment, a method for the prophylaxis and/or treatment of psoriasis.

A METHOD FOR THE PROPHYLAXIS AND/OR TREATMENT OF MEDICAL DISORDERS

FIELD OF THE INVENTION

The present invention relates generally to a method for the prophylaxis and/or treatment of medical disorders, and in particular proliferative and/or inflammatory skin disorders, and to genetic molecules useful for same. The present invention is particularly directed to genetic molecules capable of modulating growth factor interaction with its receptor on cells such as epidermal keratinocytes to inhibit, reduce or otherwise decrease stimulation of this layer of cells. The present invention contemplates, in a particularly preferred embodiment, a method for the prophylaxis and/or treatment of psoriasis or neovascularization conditions such as neovascularization of the retina. The present invention is further directed to the subject genetic molecules in adjunctive therapy for epidermal hyperplasia, such as in combination with UV treatment, and to facilitate apoptosis of cancer cells and in particular cancer cells comprising keratinocytes.

BACKGROUND OF THE INVENTION

Bibliographic details of the publications numerically referred to in this specification are collected at the end of the description.

The reference to any prior art in this specification is not, and should not be taken as, an acknowledgment or any form of suggestion that that prior art forms part of the common general knowledge in Australia or any other country.

Psoriasis and other similar conditions are common and often distressing proliferative and/or inflammatory skin disorders affecting or having the potential to affect a significant proportion of the population. The condition arises from over proliferation of basal keratinocytes in the epidermal layer of the skin associated with inflammation in the underlying dermis. Whilst a

range of treatments have been developed, none is completely effective and free of adverse side effects. Although the underlying cause of psoriasis remains elusive, there is some consensus of opinion that the condition arises at least in part from over expression of local growth factors and their interaction with their receptors supporting keratinocyte proliferation *via* keratinocyte receptors which appear to be more abundant during psoriasis.

One important group of growth factors are the dermally-derived insulin-like growth factors (IGFs) which support keratinocyte proliferation. In particular, IGF-I and IGF-II are ubiquitous peptides each with potent mitogenic effects on a broad range of cells. Molecules of the IGF type are also known as "progression factors" promoting "competent" cells through DNA synthesis. The IGFs act through a common receptor known as the Type I or IGF-I receptor, which is tyrosine kinase linked. They are synthesised in mesenchymal tissues, including the dermis, and act on adjacent cells of mesodermal, endodermal or ectodermal origin. The regulation of their synthesis involves growth hormone (GH) in the liver, but is poorly defined in most tissues [1].

Particular proteins, referred to as IGF binding proteins (IGFBPs), appear to be involved in autocrine/paracrine regulation of tissue IGF availability [2]. Six IGFBPs have so far been identified. The exact effects of the IGFBPs is not clear and observed effects *in vitro* have been inhibitory or stimulatory depending on the experimental method employed [3]. There is some evidence, however, that certain IGFBPs are involved in targeting IGF-I to its cell surface receptor.

Skin, comprising epidermis and underlying dermis, has GH receptors on dermal fibroblasts [4]. Fibroblasts synthesize IGF-I as well as IGFBPs-3, -4, -5 and -6 [5] which may be involved in targeting IGF-I to adjacent cells as well as to the overlaying epidermis. The major epidermal cell type, the keratinocyte, does not synthesize IGF-I, but possesses IGF-I receptors and is responsive to IGF-I [6].

It is apparent, therefore, that IGF-I and other growth promoting molecules, are responsible for or at least participate in a range of skin cell activities. In accordance with the present invention, the inventors have established that aberrations in the normal functioning of these molecules or

aberrations in their interaction with their receptors is an important factor in a variety of medical disorders such as proliferative and/or inflammatory skin disorders. It is proposed, therefore, to target these molecules or other molecules which facilitate their functioning or interaction with their receptors to thereby ameliorate the effects of aberrant activity during or leading to skin disease conditions and other medical conditions such as those involving neovascularization. Furthermore, these molecules may also be used to facilitate apoptosis of target cells and may be useful as adjunctive therapy for epidermal hyperplasia.

SUMMARY OF THE INVENTION

Nucleotide and amino acid sequences are referred to by a sequence identifier, i.e. (<400>1), (<400>2), etc. A sequence listing is provided after the claims.

Throughout this specification, unless the context requires otherwise, the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated element or integer or group of elements or integers but not the exclusion of any other element or integer or group of elements or integers.

Accordingly, one aspect of the present invention contemplates a method for ameliorating the effects of a medical disorder such as a proliferative and/or inflammatory skin disorder in a mammal, said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation or a cell otherwise involved in the said medical disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing a growth factor mediated cell proliferation and/or inflammation and/or other medical disorder.

According to this preferred embodiment, there is provided a method for ameliorating the effects of a medical disorder such as a proliferative and/or inflammatory skin disorder in a mammal, said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation or a cell otherwise involved with said medical disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof

capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation and/or inflammation and/or other medical disorder.

According to this embodiment, there is provided a method for ameliorating the effects of a proliferative and/or inflammatory skin disorder such as psoriasis said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation with effective amounts of UV treatment and a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation and/or inflammation.

According to this embodiment, there is provided in a particularly preferred aspect a ribozyme comprising a hybridising region and a catalytic region wherein the hybridising region is capable of hybridising to at least part of a target mRNA sequence transcribed from a genomic gene corresponding to <400>1 or <400>2 wherein said catalytic domain is capable of cleaving said target mRNA sequence to reduce or inhibit IGF-I mediated cell proliferation and/or inflammation and/or other medical disorders.

Yet another aspect of the present invention contemplates co-suppression to reduce expression or to inhibit translation of an endogenous gene encoding, for example, IGF-I, its receptor, or IGFBPs such as IGFBP-2 and/or -3. In co-suppression, a second copy of an endogenous gene or a substantially similar copy or analogue of an endogenous gene is introduced into a cell following topical administration. As with antisense molecules, nucleic acid molecules defining a ribozyme or nucleic acid molecules useful in co-suppression may first be protected such as by using a nonionic backbone.

Another aspect of the present invention contemplates a pharmaceutical composition for topical administration which comprises a nucleic acid molecule capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation such as psoriasis and one or more pharmaceutically acceptable carriers and/or diluents.

Yet another aspect of the present invention contemplates the use of a nucleic acid molecule in the manufacture of a medicament for the treatment of proliferative and/or inflammatory skin disorders or other medical disorders mediated by a growth factor.

Still a further aspect of the present invention contemplates an agent comprising a nucleic acid molecule as hereinbefore defined useful in the treatment of proliferative and/or inflammatory skin disorders, such as psoriasis or other medical disorder..

The present invention further contemplates the use of the genetic molecules and in particular the antisense molecules to inhibit the anti-apoptotic activity of IGF-I receptor.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 is a representation of the nucleotide sequence of IGFBP-2.

```
LOCUS       HSIGFBP2     1433 bp    RNA             PRI       31-JAN-1990
DEFINITION  Human mRNA for insulin-like growth factor binding protein (IGFBP-2)
ACCESSION   X16302
KEYWORDS    insulin-like growth factor binding protein.
SOURCE      human
  ORGANISM  Homo sapiens
            Eukaryota; Animalia; Metazoa; Chordata; Vertebrata; Mammalia;
            Theria; Eutheria; Primates; Haplorhini; Catarrhini; Hominidae.
REFERENCE   1  (bases 1 to 1433)
  AUTHORS   Binkert,C., Landwehr,J., Mary,J.L., Schwander,J. and Heinrich,G.
  TITLE     Cloning, sequence analysis and expression of a cDNA encoding a
            novel insulin-like growth factor binding protein (IGFBP-2)
  JOURNAL   EMBO J. 8, 2497-2502 (1989)
  STANDARD  full automatic
COMMENT     NCBI gi: 33009
FEATURES             Location/Qualifiers
     source          1. .1433
                     /organism="Homo sapiens"
                     /dev_stage="fetal"
                     /tissue_type="liver"
     misc_feature    1416. .1420
                     /note="pot. polyadenylation signal"
     polyA_site      1433
                     /note="polyadenylation site"
     CDS             118. .1104
                     /note="precursor polypeptide;  (AA -39 to 289);  NCBI gi:
                     33010."
                     /codon_start=1
                     /translation="MLPRVGCPALPLPPPPLLPLLPLLLLLLGASGGGGGARAEVLFR
                     CPPCTPERLAACGPPPVAPPAAVAAVAGGARMPCAELVREPGCGCCSVCARLEGEACG
                     VYTPRCGQGLRCYPHPGSELPLQALVMGEGTCEKRRDAEYGASPEQVADNGDDHSEGG
                     LVENHVDSTMNMLGGGGSAGRKPLKSGMKELAVFREKVTEQHRQMGKGGKHHLGLEEP
                     KKLRPPPARTPCQQELDQVLERISTMRLPDERGPLEHLYSLHIPNCDKHGLYNLKQCK
                     MSLNGQRGECWCVNPNTGKLIQGAPTIRGDPECHLFYNEQQEACGVHTQRMQ"
                     (<400>21)
     CDS             118. .234
                     /note="signal peptide;  (AA -39 to -1);  NCBI gi: 33011."
                     /codon_start=1
                     /translation="MLPRVGCPALPLPPPPLLPLLPLLLLLLGASGGGGGARA"
                     (<400>22)
     CDS             235. .1101
                     /note="mature IGFBP-2;  (AA 1 to 289);  NCBI gi: 33012."
                     /codon_start=1
                     /translation="EVLFRCPPCTPERLAACGPPPVAPPAAVAAVAGGARMPCAELVR
                     EPGCGCCSVCARLEGEACGVYTPRCGQGLRCYPHPGSELPLQALVMGEGTCEKRRDAE
                     YGASPEQVADNGDDHSEGGLVENHVDSTMNMLGGGGSAGRKPLKSGMKELAVFREKVT
                     EQHRQMGKGGKHHLGLEEPKKLRPPPARTPCQQELDQVLERISTMRLPDERGPLEHLY
                     SLHIPNCDKHGLYNLKQCKMSLNGQRGECWCVNPNTGKLIQGAPTIRGDPECHLFYNE
                     QQEACGVHTQRMQ" (<400>23)
BASE COUNT      239 a    466 c    501 g    227 t
ORIGIN
```

HSIGFBP2 Length: 1433 May 11, 1994 10:06 Type: N Check: 6232 ..

Figure 2 is a representation of the nucleotide sequence of IGFBP-3.

```
LOCUS       HUMGFIBPA    2474 bp ss-mRNA            PRI       15-JUN-1990
DEFINITION  Human growth hormone-dependent insulin-like growth factor-binding
            protein mRNA, complete cds.
ACCESSION   M31159
KEYWORDS    insulin-like growth factor binding protein.
SOURCE      Human plasma, cDNA to mRNA, clone BP-53.
  ORGANISM  Homo sapiens
            Eukaryota; Animalia; Chordata; Vertebrata; Mammalia; Theria;
            Eutheria; Primates; Haplorhini; Catarrhini; Hominidae.
REFERENCE   1  (bases 1 to 2474)
  AUTHORS   Wood,W.I., Cachianes,G., Henzel,W.J., Winslow,G.A., Spencer,S.A.,
            Hellmiss,R., Martin,J.L. and Baxter,R.C.
  TITLE     Cloning and expression of the growth hormone-dependent insulin-like
            growth factor-binding protein
  JOURNAL   Mol. Endocrinol. 2, 1176-1185 (1988)
  STANDARD  full automatic
COMMENT     NCBI gi: 183115
FEATURES             Location/Qualifiers
     mRNA            <1. .2474
                     /note="GFIBP mRNA"
     CDS             110. .985
                     /gene="IGFBP1"
                     /note="insulin-like growth factor-binding protein;  NCBI
                     gi: 183116."
                     /codon_start=1
                     /translation="MQRARPTLWAAALTLLVLLRGPPVARAGASSGGLGPVVRCEPCD
                     ARALAQCAPPPAVCAELVREPGCGCCLTCALSEGQPCGIYTERCGSGLRCQPSPDEAR
                     PLQALLDGRGLCVNASAVSRLRAYLLPAPPAPGNASESEEDRSAGSVESPSVSSTHRV
                     SDPKFHPLHSKIIIIKKGHAKDSQRYKVDYESQSTDTQNFSSESKRETEYGPCRREME
                     DTLNHLKFLNVLSPRGVHIPNCDKKGFYKKKQCRPSKGRKRGFCWCVDKYGQPLPGYT
                     TKGKEDVHCYSMQSK" (<400>24>)
     source          1. .2474
                     /organism="Homo sapiens"
BASE COUNT      597 a    646 c    651 g    580 t
ORIGIN
```

HUMGFIBPA Length: 2474 May 11, 1994 10:00 Type: N Check: 9946 ..

Figure 3 is a representation of the nucleotide sequence of IGF-1-receptor.

```
LOCUS       HSIGFIRR     4989 bp    RNA             PRI       28-MAR-1991
DEFINITION  Human mRNA for insulin-like growth factor I receptor
ACCESSION   X04434 M24599
KEYWORDS    glycoprotein; insulin receptor;
            insulin-like growth factor I receptor; membrane glycoprotein;
            receptor; tyrosine kinase.
SOURCE      human
```

```
     ORGANISM  Homo sapiens
               Eukaryota; Animalia; Metazoa; Chordata; Vertebrata; Mammalia;
               Theria; Eutheria; Primates; Haplorhini; Catarrhini; Hominidae.
  REFERENCE    1  (bases 1 to 4989)
    AUTHORS    Ullrich,A., Gray,A., Tam,A.W., Yang-Feng,T., Tsubokawa,M.,
               Collins,C., Henzel,W., Bon,T.L., Kathuria,S., Chen,E., Jakobs,S.,
               Francke,U., Ramachandran,J. and Fujita-Yamaguchi,Y.
    TITLE      Insulin-like growth factor I receptor primary structure: comparison
               with insulin receptor suggests structural dererminants that define
               functional specificity
    JOURNAL    EMBO J. 5, 2503-2512 (1986)
    STANDARD   full automatic
  COMMENT      NCBI gi: 33058
  FEATURES               Location/Qualifiers
       source            1. .4989
                         /organism="Homo sapiens"
                         /tissue_type="placenta"
                         /clone_lib="(lamda)gt10"
                         /clone="(lambda)IGF-1-R.85, (lambda)IGF-1-R.76"
       sig_peptide       32. .121
       mat_peptide       122. .4132
                         /note="IGF-I receptor"
       misc_feature      122. .2251
                         /note="alpha-subunit (AA 1 - 710)"
       misc_feature      182. .190
                         /note="pot.N-linked glycosylation site (AA 21 - 23)"
       misc_feature      335. .343
                         /note="pot.N-linked glycostlation site (AA 72 - 74)"
       misc_feature      434. .442
                         /note="pot.N-linked glycostlation site (AA 105 - 107)"
       misc_feature      761. .769
                         /note="pot.N-linked glycostlation site (AA 214 - 216)"
       misc_feature      971. .979
                         /note="pot.N-linked glycostlation site (AA 284 - 286)"
       misc_feature      1280. .1288
                         /note="pot.N-linked glycostlation site (AA 387 - 389)"
       misc_feature      1343. .1351
                         /note="pot.N-linked glycosylation site (AA 408 - 410)"
       misc_feature      1631. .1639
                         /note="pot.N-linked glycostlation site (AA 504 - 506)"
       misc_feature      1850. .1858
                         /note="pot.N-linked glycosylation site (AA 577 - 579)"
       misc_feature      1895. .1903
                         /note="pot.N-linked glycosylation site (AA 592 - 594)"
       misc_feature      1949. .1957
                         /note="pot.N-linked glycosylation site (AA 610 - 612)"
       misc_feature      2240. .2251
                         /note="putative proreceptor processing site (AA 707 -
                         710)"
       misc_feature      2252. .4132
                         /note="beta-subunit (AA 711 - 1337)"
       misc_feature      2270. .2278
                         /note="pot.N-linked glycosylation site (AA 717 - 719]"
       misc_feature      2297. .2305
                         /note="pot.N-linked glycosylation site (AA 726 - 728)"
       misc_feature      2321. .2329
                         /note="pot.N-linked glycosylation site (AA 734 - 736)"
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     misc_feature    2729. .2737
                     /note="pot.N-linked glycosylation site (AA 870 - 872)"
     misc_feature    2768. .2776
                     /note="pot.N-linked glycosylation site (AA 883 - 885)"
     misc_feature    2837. .2908
                     /note="transmembrane region (AA 906 - 929)"
     misc_feature    2918. .2926
                     /note="pot.N-linked glycosylation site (AA 933 - 935)"
     misc_feature    3047. .3049
                     /note="pot.ATP binding site (AA 976)"
     misc_feature    3053. .3055
                     /note="pot.ATP binding site (AA 978)"
     misc_feature    3062. .3064
                     /note="pot.ATP binding site (AA 981)"
     misc_feature    3128. .3130
                     /note="pot.ATP binding site (AA 1003)"
     CDS             32. .4132
                     /product="IGF-I receptor"
                     /note="50 stops when translation attempted, frame 1, code
                     0"
BASE COUNT     1216 a   1371 c   1320 g   1082 t
ORIGIN

  HSIGFIRR  Length: 4989  May 11, 1994  12:10  Type: N  Check: 133  ..
```

Figure 4A is a photographic representation of a Western ligand blot of HaCaT conditioned medium showing IGFBP-3 secreted in 24 hours after 7 day treatment with phosphorothioate oligonucleotides (BP3AS2, BP3AS3 and BP3S) at 0.5μM and 5μM;

* no oligonucleotide added.

Figure 4B is a graphical representation of a scanning imaging desitometry of Western ligand blot (Figure 4A), showing relative band intensities of IGFBP-3 and the 24kDa IGFBP-4 after treatment with phosphorothioate oligonucleotides;

* no oligonucleotide added.

Figure 5A is a photographic representation of a Western ligand blot of HaCaT conditioned medium showing IGFBP-3 secreted in 24 hours after 7 day treatment with phosophorothioate oligonucleotide BP3AS2 at 0.5μM compared with several control oligonucleotides at 0.5μM. (a) oligonucleotide BP3AS2NS; (b) oligonucleotide BP3AS4; (c) oligonucleotide BP3AS4NS; and (untreated), no oligonucleotide added.

Figure 5B is a graphical representation of a scanning imaging densitometry of Western ligand blot (Figure 5A), showing relative band intensities of IGFBP-3 after treatment with phosphorothioate oligonucleotides as in Figure 5A, showing IGFBP-3 band intensities expressed as a percentage of the average band intensity from conditioned medium of cells not treated with oligonucleotide.

Figure 6 is a graphical representation showing inhibition of IGF-I binding by antisense oligonucleotides to IGF-I receptor. IGFR.AS: antisense; IGFR.S: sense.

Figure 7 is a graphical representation showing inhibition of IGFBP-3 production in culture medium following initial treatment with antisense oligonucleotides once daily over a 2 day period.

Figure 8 is a graphical representation showing optimization of IGFBP-3 antisense oligonucleotide concentration as determined by relative IGFBP-3 concentration in culture medium.

Figure 9 is a diagramatic representation of a map of IGF-1 Receptor mRNA and position of target ODNs.

Figure 10 is a photographical representation showing Lipid-mediated uptake of oligonucleotide in keratinocytes. HaCaT keratinocytes were incubated for 24 hours in medium (DMEM plus 10% v/v FCS) containing fluorescently labelled ODN (R451, 30 nM) and cytofectin GSV (2 μg/ml). The cells were then transferred to ODN-free medium and fluorescence microscopy (a) and phase contrast (b) images of the cells were obtained.

Figure 11 is a graphical representation of uptake (A) and toxicity (B) of ODN/lipid complexes in keratinocytes. Confluence HaCaT keratinocytes were incubated in DMEM containing fluoresently labelled ODN (R451) plus liposome over 120 hours, viewed using fluorescene microscopy and trypan blue stained and counted.

Figure 12 is a graphical representation of an IGF-1 Receptor mRNA in ODN treated (30nM) HaCaT cells (2μg/ml GSV). HaCaT keratinocytes were treated for 96 hours with C-5 propynyl, dU, dC ODNs complexed with cytofectin GSV. Cells were treated with ODNs complementary to the human IGF-I receptor mRNA (27, 32, 74 and 78), 2 randomised sequence ODNs (R451) and R766), liposome alone (GSV) or were left untreated (UT). Total RNA was isolated then analysed for IGF-I receptor mRNA and GAPDH mRNA levels by RNase Protection and PhosphorImager quantitiation.

(A) Electrophoretic analysis of IGF-I receptor and GAPDH mRNA fragments after RNase Protection. Molecular weight markers are shown on the right hand side. Full length probe is shown on the left hand side (G-probe and I-probe). GAPDH protected fragments (G) are seen at 316 bases and IGF-I receptor protected fragments (I) are seen at 276 bases.

(B) Relative level of IGF-I receptor mRNA following each treatment is shown.

Figure 13 is a graphical representation of an IGF-1 receptor mRNA in ODN treated (30nM) HaCaT cells (2μg/ml GSV). Summary of IGF-I receptor ODN screening data. HaCaT keratinocytes were treated for 96 hours with C-5 propynyl, dU, dC ODNs complexed with cytofectin GSV. Total RNA was isolated then analysed for IGF-I receptor mRNA and GAPDH mRNA levels by RNase protection and phosphorImager quantitiation. Relative level of IGF-I receptor mRNA is shown after treatment with ODNs complementary to the human IGF-I receptor mRNA, 4 randomised sequence ODNs and liposome alone. (26-86=IGF-I receptor ODNs; R1, R4, R7 and R9 = randomised ODNs (R1=R121, R4=R451, R7=R766, R9=R961); GSV=liposome alone; UT=untreated). *indicates a significant difference in relative IGF-I receptor mRNA from GSV treated cells (n=4-10, $p<0.05$).

Figure 14 is a graphical representation of the effect of antisense oligonucleotides on IGF-1 receptor levels on the surface of keratinocytes. HaCaT cells were grown to confluence in 24-well plates in DMEM containing 10% v/v FCS. Oligodeoxynucleotide (ODN) and Cytofectin GSV (GSV, Glen Research) were mixed together in serum-free DMEM, incubated at room

temperature for 10 minutes before being diluted ten-fold in medium and placed on the cells. Cells were incubated for 72 hours with 30 nM random sequence or antisense ODN and 2 μg/ml GSV or with GSV alone in DMEM containing 10% v/v FCS with solutions replaced every 24 hours. This was followed by incubation with ODN/GSV in serum-free DMEM for 48 hours. All incubations were performed at 37°C. Wells were washed twice with 1 ml cold PBS. Serum-free DMEM containing 10^{-10}M ^{125}I-IGF-I was added with or without the IGF-I analogue, des (1-3) IGF-I, at 10^{-10}M to 10^{-7}M. Cells were incubated at 4°C for 17 hours with gentle shaking then washed three times with 1 ml cold PBS and lysed in 250 μl 0.5M NaOH/0.1% v/v Triton X-100 at room temperature for 4 hours. Specific binding of the solubilised cell extract was measured using a γ counter.

Figure 15 is a graphical representation of the effect of antisense oligonucleotides on IGF-1 receptor levels on the surface of keratinocytes.

Figure 16 is a photographical representation of H & E stained sections of (A) psoriatic skin biopsy prior to grafting and (B) 49 day old psoriatic skin graft using skin from the same donor.

Figure 17 is a photographical representation of uptake of oligonucleotide after intradermal injection into psoriatic skin graft on a nude mouse. Psoriatic skin graft was intradermally injected with ODN (R451, 50 μl, 10 μM). The graft was removed and sectioned after 24 hours, then viewed using confocal microscopy.

Figure 18(a) is a photographical representation of Pregraft, Donor JH, Donor JH, PBS treated, 50μl, Donor JH, #50 treated, 50μl, 10μM.

Figure 18(b) is a photographical representation of Donor LB, pregraft, Donor LB, PBS treated (50μl), Donor LB, #74 treated (50μl, 10μM).

Figure 18(c) is a photographical representation of Donor PW, pregraft, Donor PW, R451 treated (50μl, 10μM), Donor LB, #74 treated (50μl, 10μM).

Figure 18(d) is a photographical representation of Donor GM, pregraft, Donor GB, R451 treated (50μl, 10μM), Donor GM, #27 treated (50μl, 10μM).

Figure 19(a) is a photographical representation showing Donor JH pregraft, Donor JH PBS treated 50μl, Donor JH #50 treated 50μl, 10μM.

Figure 19(b) is a photographical representation Donor LB pregraft, Donor LB PBS treated 50μl, Donor LB #74 treated 50μl, 10μM.

Figure 19(c) is a photographical representational showing Donor PW pregraft, Donor PW r451 treated 50μl, 10μM, Donor PW #74 treated 50μl, 10μM.

Figure 19(d) is a photographical representation showing Donor GM pregraft, Donor GM R451 treated 50μl, 10μM, Donor #27 treated 50μl, 10μM.

Figure 20 is a graphical representation showing suppression of psoriasis after treatment with oligonucleotide (quantification). Oligonucleotide (50 μl, 10μM) was injected every two days for 20 days, as were control treatments. Skin thickness was measured by removing the skin and using computer software (MCID analysis) to measure the exact thickness of each graft. N=3-4 for each treatment. *indicates a significant difference from the pregraft value (ANOVA, $P<0.05$)

Figure 21 is a photographic representation of αhKi-67 imunobiological binding.

Figure 22 is a photographical representation showing penetration of oligonucleotide into human skin after topical treatment. Fluorescently labelled oligonucleotide (10 μM R451) was

applied topically after formulation with cytofectin GSV (10 μg/ml) and viewed using confocal microscopy.

Figure 23 is a photographical representation showing penetration of oligonucleotide into human skin after application of topical gel formation. Fluorescently labelled oligonucleotide (10 μM R451) was applied topically after complexing with cytofectin GSV (10 μg/ml) and formulation into 3% methylcellulose gel. Image was obtained using confocal microscopy.

Figure 24 is a graphical representation showing IGFBP-3 mRNA.

Figure 25(a) is a graphical representation showing IGFBP-3 mRNA in AON treated (100nM) HaCaT cells (2μg/ml GSV).

Figure 25(b) is a graphical representation showing IGFBP-3 mRNA levels of AON treated (100nm) HaCaT cells (2μg/ml GSV).

Figure 25(c) is a graphical representation showing IGFBP-3 mRNA in AON treated (30nM) HaCaT cells (2μg/ml GSV).

Figure 25(d) is a graphical representation showing IGFBP-3 mRNA in AON treated (30nM) HaCaT cells (2μg/ml GSV).

Figure 26(a) is a graphical representation showing IGFBP-3 mRNA in ODN treated (30nM) HaCaT cells (2μg/ml). HaCaT keratinocytes were treated for 51 hours with C-5 propynl, dU, dC ODNs complexed with cytofectin GSV. Total RNA was isolated then analysed for IGFBP-3 mRNA and GAPDH mRNA levels by Northern analysis and phosphorimager quantitation. Relative level of IGFBP-3 mRNA is shown after treatment with ODNs complementary to the human IGFBP-3 mRNA, 4 randomised sequence ODNs and lipsome alone. (1-24=IGFBP-3 ODNs; R1, R4, R7 and R9=randomised ODNs (R1=R121, R4=R451, R7=R766, R9 R961); GS=liposome alone; UT=untreated). *indicates a significant different in relative

IGFBP-3 mRNA from GSV treated cells (n- 5-8, $p<0.01$), **indicates a significant difference in relative IGFBP-3 mRNA from GSV treated cells (n= 5-8, $p<0.05$).

Figure 26(b) is a graphical representation showing IGFBP-3 mRNA in ODN treated (100nM) HaCaT cells (2μg/ml GSV). HaCaT keratinocytes were treated for 51 hours with C-5 propynl, dU, dC ODNs complexed with cytofectin GSV. Total RNA was isolated then analysed for IGFBP-3 mRNA and GAPDH mRNA levels by Northern analysis and phosphorimager quantitation. Relative level of IGFBP-3 mRNA is shown after treatment with ODNs complementary to the human IGFBP-3 mRNA, 4 randomised sequence ODNs and liposome alone. (1-24=IGFBP-3 ODNs; R1, R4, R7 and R9 = randomised ODNs (R1-R121, R4=R451, R7=R766, R9-R961), GS=lipsome alone; UT=untreated). *indicates a significant difference in relative IGFBP-3 mRNA from GSV treated cells (n- 6-8, $p<0.01$).

Figure 27 is a representation showing a reduction in IGF-I receptor mRNA in HaCaT cells following treatment with antisense oligonucleotides. Confluent HaCaT cells were treated every 24 h for 4 days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM IGF-I receptor specific oligonucleotides *(#26 to #86)* or random sequence oligonucleotides (R121, R451 and R766). Total RNA was isolated and analysed for IGF-I receptor and GAPDH mRNA by RNase protection assay. (a). Representative RNase protection assay gel showing IGF-I receptor *(IGF*R) and GAPDH mRNA in untreated or treated HaCaT cells. In this example, a reduction in IGFR band intensity relative to GAPDH can be seen with AON #27 and #78, but not with #32, #74 or the controls *(R*4, *R*7, random oligonucleotides R451 and R766, respectively; G, GSV lipid; *U*T, untreated).

(b) Densitometric quantitation of IGF-I receptor mRNA (normalised to GAPDH mRNA) in HaCaT cells following treatment with IGF-I receptor specific oligonucleotides (solid black), random sequence oligonucleotides (horizontal striped bar) or GSV alone (shaded bar) compared to untreated cells *(U*T, vertical striped bar). Each oligonucleotide was assayed in duplicate in at least two separate experiments.

Results are presented as mean ± SEM. A one-way ANOVA followed by Tukey's (▲) test was performed; ▲ indicates a significant difference between cells treated with IGF-I receptor specific AONs and all of the control treatments ($p<0.05$). n=4 except for #27 and #32 (n=6), #28 and #68 (n=3), R766 (n=9), and R451, GSV and untreated (n=10).

Figure 28 is a representation showing a reduction in total cellular IGF-I receptor protein following antisense oligonucleotide treatment. Confluent HaCaT cells were treated every 24 h for 4 days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM IGF-I receptor specific AONs (#27, #50 and #64) or the random sequence oligonucleotide, R451. Total cellular protein was isolated and analysed for IGF-I receptor by SDS PAGE followed by western blotting with an antibody specific for the human IGF-I receptor. **(a)** Duplicate treated cellular extracts showing the IGF-I receptor at the predicted size of 110 kD

(b) Densitometric quantitation of IGF-I receptor protein. Results are presented as mean ± SEM of four different experiments each performed in duplicate. A one-way ANOVA followed by a Dunnett's test was performed; * indicates a significant difference from GSV treated cells ($p<0.01$). *GSV*, GSV lipid alone; *UT*, untreated; R451, random sequence oligonucleotide; *64*, *50*, *27*, IGF-I receptor-specific AONs.

Figure 29 is a representation showing a reduction in IGF-I receptor numbers on the keratinocyte cell surface after antisense oligonucleotide treatment. HaCaT cells were transfected with IGF-I receptor specific AONs #27 (—▲—), #50 (—x—), #64 (---■---), a random sequence oligonucleotide R451 (—o—), or treated with GSV lipid alone (--□--) every 24 h for four days (untreated cells, --✳--). Competition binding assays using ^{125}I-IGF-I and the receptor-specific analogue, des(1-3)IGF-I, were performed (inset); plotted values are means ± standard error. The mean values were then subjected to Scatchard analysis.

Figure 30 is a representation showing a reduction in keratinocyte cell number following antisense oligonucleotide treatment. HaCaT cells, initially at 40% confluence, were transfected with the IGF-I receptor specific AON #64, control sequences R451 and 6416, or

treated with GSV lipid alone every 24 h for 2 days (UT, untreated cells). Cell number was measured in the culture wells using a dye binding assay (Experimental protocol). Results are presented as mean ± SD. A one-way ANOVA was performed, followed by a Tukey's multiple comparison test. ▲ indicates a significant difference between cells treated with AON #64 and all of the control treatments ($p<0.001$).

Figure 31 is a representation showing a reversal of epidermal hyperplasia in psoriatic human skin grafts on nude mice following intradermal injection with antisense oligonucleotides

Grafted psoriasis lesions were injected with IGF-I receptor specific AONs, a random sequence oligonucleotide in PBS, or with PBS alone, every 2 days for 20 days, then analysed histologically. **(a)** Donor A graft treated with AON #50 showing epidermal thinning compared with pregraft and control *(PBS)* treated graft, and Donor B graft treated with AON #27 showing epidermal thinning compared with pregraft and control (R451) treated graft. E, epidermis; *Scale b*ar, 400 mm; all pictures are at the same magnification. **(b)** Mean epidermal cross-sectional area over the full width of grafts was determined by digital image analysis. Results are presented as mean ± SEM. *Shaded bar*s, control treatments: R451, random oligonucleotide sequence; *solid bar*s, treatments with oligonucleotides that inhibited IGF-I receptor expression in vitro. * indicates a significant difference from the vehicle treated graft ($p<0.01$, n=5-7), ++ indicates a significant difference from the random sequence (R451) treated graft ($p<0.01$, n=5-7). **(c)** Parakeratosis *(arrow)* was absent in grafts treated with IGF-I receptor AONs *(AON #50)* but persisted in pregraft and control *(PBS)* treated graft. *Scale bar, 100 mm.*

Figure 32 is a representation showing a reversal of epidermal hyperplasia correlates with reduced IGF-I receptor mRNA in grafted psoriasis lesions treated with antisense oligonucleotides (a) A psoriasis lesion prior to grafting, and after grafting and treatment with IGF-I receptor specific oligonucleotide #27 *(AON #27)* or random sequence *(R451)* was immunostained with antibodies to Ki67 to identify proliferating cells. Proliferating cells are indicated by a dark brown nucleus (arrows). *Scale bar, 250 m*m; all pictures are at the same

magnification. **(b)** The same lesion prior to grafting and after oligonucleotide treatment as in (a) was subjected to in situ hybridisation with a ^{35}S-labeled cRNA probe complementary to the human IGF-I receptor mRNA. The presence of IGF-I receptor mRNA is indicated by silver grains (tiny black speckles), which are almost eliminated in the epidermis of the lesion treated with the IGF-I receptor-specific oligonucleotide #27 (AON #27). Arrows indicate the basal layer of the epidermis with dermis underneath. *Scale bar, 50 μm.*

Figure 33 is a representation showing a reduction in IGF-I receptor mRNA in HaCaT keratinocytes following treatment with oligonucleotides. HaCaT cell monolayers grown to 90% confluence in DMEM contianing 10% v/v fetal calf serum were treated with 24 h for two days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM oligonucleotide. Total RNA was isolated and analysed for IGF-I receptor and GAPDH mRNA using a commercially availble ribonuclease protection assay kit (RPAII, Ambicon Inc, Austin, Texas). Band intensity was quantified using ImageQuant software (Molecular Dynamics, Sunnyvale, California).

Figure 34 is a representation showing a reduction in IGF-I receptor protein in HaCaT keratinocytes following treatment with oligonucleotides. HaCaT cell monolayers grown to 90% confluence in DMEM containing 10% v/v fetal calf serum were treated every 24 h for four days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM oligonucleotide. Cells were lyased in a buffer containing 50 mM HEPES, 150 mM NaCl, 10% v/v gycerol, 1% v/v Triton X-100 and 100 μg/ml aprotinin on ice for 30 mins, then 30 μg of lysate was loaded onto a denaturing 7% w/v polyacrylamide gel followed by transfer onto an Immobilon-P membrane (Millipore, Bedford, Massachusetts). Membranes were incubated with the anti-IGF-I receptor antibody C20 (Sanra Cruz Biotechnology Inc., Santa Cruz, California, 25 ng/ml in 150 mM NaCl, 10 mM Tris-HCl, pH 7.4, 0.1% v/v Tween 20) for 1 h at room temperature and developed using the Vistra ECF western blotting kit (Amersham, Buckinghamshire, England). Band intensity was quantified using ImageQuant software (Molecular Dynamics, Sunnyvale, California).

Figure 35 is a representation showing a reduction in HaCaT keratinocyte cell number following treatment with oligonucleotides. HaCaT cell monolayers grown to 40% confluence in DMEM containing 10% fetal calf serum were treated every 24 h for three days with 2 μg/ml GSV lipid alone (GSV) or complexed with 15 nM oligonucleotide. Cell number was measured every 24 h using the amido black dye binding assay [32].

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

The present invention is predicated in part on the use of molecules and in particular genetic molecules and more particularly antisense molecules to down-regulate a growth factor, its receptor and/or growth factor expression facilitating sequences.

Accordingly, one aspect of the present invention contemplates a method for ameliorating the effects of a medical disorder such as a proliferative and/or inflammatory skin disorder in a mammal, said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation or a cell otherwise involved in the said medical disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing a growth factor mediated cell proliferation and/or inflammation and/or other medical disorder.

Growth factor mediated cell proliferation and inflammation are also referred to as epidermal hyperplasias and these and other medical disorders may be mediated by any number of molecules such as but not limited to IGF-I, keratinocyte growth factor (KGF), transforming growth factor-α (TGFα), tumour necrosis factor-α (TNFα), interleukin-1, -4, -6 and 8 (IL-1, IL-4, IL-6 and IL-8, respectively), basic fibroblast growth factor (bFGF) or a combination of one or more of the above. The present invention is particularly described and exemplified with reference to IGF-I and its receptor (IGF-I receptor) and to IGF-I facilitating molecules, IGFBPs, since targeting these molecules according to the methods contemplated herein provides the best results to date. This is done, however, with the understanding that the present invention extends to any growth factor or cytokine-like molecule, a receptor thereof or a facilitating molecule like the IGFBPs involved in skin cell proliferation such as those molecules contemplated above and/or their receptors and/or facilitating molecules therefor.

According to this preferred embodiment, there is provided a method for ameliorating the effects of a medical disorder such as a proliferative and/or inflammatory skin disorder in a mammal, said method comprising contacting the proliferating and/or inflamed skin or skin

capable of proliferation and/or inflammation or a cell otherwise involved with said medical disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation and/or inflammation and/or other medical disorder.

The present invention is particularly described by psoriasis as the proliferative skin disorder. However, the subject invention extends to a range of proliferative and/or inflammatory skin disorders or epidermal hyperplasias such as but not limited to psoriasis, ichthyosis, pityriasis rubra pilaris ("PRP"), seborrhoea, keloids, keratoses, neoplasias and scleroderma, warts, benign growths and cancers of the skin. The present invention extends to a range of other disorders such as neovascularization conditions such as but not limited to hyperneovasularization such as neovascularization of the retina, lining of the brain, skin, hyperproliferation of the inside of blood vessels, kidney disease, atherosclerotic disease, hyperplasias of the gut epithelium or growth factor mediated malignancies such as IGF1-mediated malignancies.

Furthermore, down-regulation of IGF-I receptor is useful as adjunctive therapy for epidermal hyperplasia. In accordance with this aspect of the present invention it is known that IGF-I receptor elicits separate intracellular signals which prevent apoptosis [19]. In keratinocytes, IGF-I receptor activation has been shown to protect UV-irradiated cells from apoptosis [20]. In another cell type, a number of IGF-I receptors expressed by the cells correlated with tumorigenicity and apoptotic resistance [21]. Consequently, in accordance with the present invention, by inactivating IGF-I receptor on cells such as epidermal keratinocytes will achieve three important outcomes:

(i) Acute epidermal hyperplasia following UV has been suggested to increase the risk of keratinocyte carcinogenic transformation [22]. By reducing IGF-I receptor expression in the epidermis, the incidence of epidermal hyperplasia following UV exposure is likely to be reduced leading to an overall acceleration in normalization of the lesion and reduced carcinogenic risk.

(ii) Inhibition of anti-apoptotic action of IGF-I receptor will enhance the reversal of epidermal thickening and accelerate normalization of differentiation. Topical or injected IGF-I receptor antisense as adjunctive treatment will increase apoptosis in the epidermal layer thereby enhancing the reduction in acanthosis observed in UV treatments.

(iii) Survival of keratinocytes, ie. those which evade apoptosis is likely to occur when cells have damaged DNA. Such mutations may be in the tumor suppressor region. Consequently, the use of antisense therapy will result in less frequent selection of mutated keratinocytes and therefore reduced incidence of basal cell carcinomas and squamous.

According to this embodiment, there is provided a method for ameliorating the effects of a proliferative and/or inflammatory skin disorder such as psoriasis said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation with effective amounts of UV treatment and a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation and/or inflammation.

The UV treatment and nucleic acid molecule or its chemical analogue may be administered in any order or may be done simultaneously. This method is particularly useful in treating psoriasis by combination of UV and antisense therapy. Preferably the antisense therapy is directed to the IGF-I receptor.

In a preferred embodiment, the present invention is directed to a method for ameliorating the effects of psoriasis or other medical disorder, said method comprising contacting proliferating skin or skin capable of proliferation or cells associated with said disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation or ameliorating the medical disorder.

The present invention extends to any mammal such as but not limited to humans, livestock animals (e.g. horses, sheep, cows, goats, pigs, donkeys), laboratory test animals (e.g. rabbits, mice, guinea pigs), companion animals (e.g. cats, dogs) and captive wild animals. However, the instant invention is particularly directed to proliferative and/or inflammatory skin disorders such as psoriasis in humans as well as medical disorders contemplated above.

The aspects of the subject invention instantly contemplated are particularly directed to the topical application of one or more suitable nucleic molecules capable of inhibiting, reducing or otherwise interfering with IGF-mediated cell proliferation and/or inflammation. More particularly, the nucleic acid molecule targets IGF-I interaction with its receptor. Conveniently, therefore, the nucleic acid molecule is an antagonist of IGF-I interaction with its receptor. Most conveniently, the nucleic acid molecule antagonist is an antisense molecule to the IGF-I receptor, to IGF-I itself or to a molecule capable of facilitating IGF-I interaction with its receptor such as but not limited to an IGFBP.

Insofar as the invention relates to IGFBPs, the preferred molecules are IGFBP-2, -3, -4, -5 and -6. The most preferred molecules are IGFBP-2 and IGFBP-3.

The nucleotide sequences of IGFBP-2 and IGFBP-3 are set forth in Figures 1 (<400>1) and 2 (<400>2), respectively. According to a particularly preferred aspect of the present invention, there is provided a nucleic acid molecule comprising at least about ten nucleotides capable of hybridising to, forming a heteroduplex or otherwise interacting with an mRNA molecule directed from a gene corresponding to a genomic form of <400>1 and/or <400>2 and which thereby reduces or inhibits translation of said mRNA molecule. Preferably, the nucleic acid molecule is at least about 15 nucleotides in length and more preferably at least about 20-25 nucleotides in length. However, the instant invention extends to any length nucleic acid molecule including a molecule of 100-200 nucleotides in length to correspond to the full length of or near full length of the subject genes.

The nucleotide sequence of the antisense molecules may correspond exactly to a region or portion of <400>1 or <400>2 or may differ by one or more nucleotide substitutions, deletions and/or additions. It is a requirement, however, that the nucleic acid molecule interact with an mRNA molecule to thereby reduce its translation into active protein.

Examples of potential antisense molecules for IGFBP-2 and IGFBP-3 are those capable of interacting with sequences selected from the lists in Examples 6 and 7, respectively.

The nucleic acid molecules in the form of an antisense molecule may be linear or covalently closed circular and single stranded or partially double stranded. A double stranded molecule may form a triplex with target mRNA or a target gene. The molecule may also be protected from, for example, nucleases, by any number of means such as using a nonionic backbone or a phosphorothioate linkage. A convenient nonionic backbone contemplated herein is ethylphosphotriester linkage or a 2′-O-methylribosyl derivative. A particularly useful modification modifies the DNA backbone by introducing phosphorothioate internucleotide linkages. Alternatively or in addition to the pyrimidine bases are modified by inclusion of a C-5 propyne substitution which modification is proposed to enhance duplex stability [23]. The present invention extends to any chemical modification to the bases and/or RNA or DNA backbone. Reference to a "chemical analogue" of a nucleic acid molecule includes reference to a modified base, nucleotide, nucleoside or phosphate backbone.

Examples of suitable oligonucleotide analogues are conveniently described in Ts'O *et al* [7]. Further suitable examples of oligonucleotide analogues and chemical modifications are described in references 25 to 31.

Alternatively, the antisense molecules of the present invention may target the IGF-I gene itself or its receptor or a multivalent antisense molecule may be constructed or separate molecules administered which target at least two or an IGFBP, IGF-I and/or IGF-I-receptor. Examples of suitable antisense molecules capable of targetting the IGF-I receptor are those capable of

interacting with sequences selected from the list in Example 8. One particularly useful antisense molecule is 5'- ATCTCTCCGCTTCCTTTC -3' (<400>10).

Other particularly useful antisense molecules are:

#27 UCCGGAGCCAGACUU

#64 CACAGUUGCUGCAAG

#78 UCUCCGCUUCCUUUC

#28 AGCCCCCACAGCGAG

#32 GCCUUGGAGAUGAGC

#40 UAACAGAGGUCAGCA

#42 GGAUCAGGGACCAGU

#46 CGGCAAGCUACACAG

#50 GGCAGGCAGGCACAC

Particularly useful molecules are selected from #27, #64 and #78. In a preferred embodiment these molecules comprise a C-5 propynyl dU, dC phosphorothioate modification.

A particularly preferred embodiment of the present invention contemplates a method of ameliorating the effects of psoriasis or other medical disorder, said method comprising contacting proliferating skin or skin capable of proliferation or cells associated with said medical disorder with an effective amount of one or more nucleic acid molecules or chemical analogues thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation or ameliorating the medical disorder wherein said one or more molecules comprises a polynucleotide capable of interacting with mRNA directed from an IGF-I gene, an IGF-I receptor gene or a gene encoding an IGFBP such as IGFBP-2 and/or IGFBP-3.

Preferably, the nucleic acid molecule are antisense molecules. Particularly useful antisense molecules are:

#27 UCCGGAGCCAGACUU

#64 CACAGUUGCUGCAAG

#78 UCUCCGCUUCCUUUC

#28 AGCCCCCACAGCGAG

#32 GCCUUGGAGAUGAGC

#40 UAACAGAGGUCAGCA

#42 GGAUCAGGGACCAGU

#46 CGGCAAGCUACACAG

#50 GGCAGGCAGGCACAC

Even more particularly useful molecules are selected from #27, #64 and #78.

In accordance with one aspect of the present invention the nucleic acid molecule is topically applied in aqueous solution or in conjunction with a cream, ointment, oil or other suitable carrier and/or diluent. A single application may be sufficient depending on the severity or exigencies of the condition although more commonly, multiple applications are required ranging from hourly, multi-hourly, daily, multi-daily, weekly or monthly, or in some other suitable time interval. The treatment might comprise solely the application of the nucleic acid molecule or this may be applied in conjunction with other treatments for the skin proliferation and/or inflammatory disorder being treated or for other associated conditions including microbial infection, bleeding and the formation of a variety of rashes.

As an alternative to or in conjunction with antisense therapy, the subject invention extends to the nucleic acid molecule as, or incorporating, a ribozyme including a minizyme to, for example, IGF-I, its receptor or to molecules such as IGFBPs and in particular IGFBP-2 and -3. Ribozymes are synthetic nucleic acid molecules which possess highly specific endoribonuclease activity. In particular, they comprise a hybridising region which is complementary in nucleotide sequence to at least part of a target RNA. Ribozymes are well described by Haseloff and Gerlach [8] and in International Patent Application No. WO 89/05852. The present invention extends to ribozymes which target mRNA specified by genes encoding IGF-I, its receptor or one or more IGFBPs such as IGFBP-2 and/or IGFBP-3.

According to this embodiment, there is provided in a particularly preferred aspect a ribozyme comprising a hybridising region and a catalytic region wherein the hybridising region is capable of hybridising to at least part of a target mRNA sequence transcribed from a genomic gene corresponding to (<400>1) or (<400>2) wherein said catalytic domain is capable of cleaving said target mRNA sequence to reduce or inhibit IGF-I mediated cell proliferation and/or inflammation and/or other medical disorders.

Yet another aspect of the present invention contemplates co-suppression to reduce expression or to inhibit translation of an endogenous gene encoding, for example, IGF-I, its receptor, or IGFBPs such as IGFBP-2 and/or -3. In co-suppression, a second copy of an endogenous gene or a substantially similar copy or analogue of an endogenous gene is introduced into a cell following topical administration. As with antisense molecules, nucleic acid molecules defining a ribozyme or nucleic acid molecules useful in co-suppression may first be protected such as by using a nonionic backbone.

The efficacy of the nucleic acid molecules of the present invention can be conveniently tested and screened using an *in vitro* system comprising a basal keratinocyte cell line. A particularly useful system comprises the HaCaT cell line described by Boukamp *et al* [9]. In one assay, IGF-I is added to an oligonucleotide treated HaCaT cell line. Alternatively, growth of oligonucleotide treated HaCaT cells is observed on a feeder layer of irradiated 3T3 fibroblasts. Using such *in vitro* assays, it is observed that antisense oligonucleotides to IGFBP-3, for example, inhibit production of IGFBP-3 by HaCaT cells. Other suitable animal models include the nude mouse/human skin graft model (15; 16) and the "flaky skin" mouse model (17; 18). In the nude mouse model, microdermatome biopsies of psoriasis lesions are taken under local anaesthetic from volunteers then transplanted to congenital athymic (nude) mice. These transplanted human skin grafts maintain the characteristic hyperproliferating epidermis for 6-8 weeks. They are an established model for testing the efficacy of topically applied therapies for psoriasis. In the "flaky skin" mouse model, the fsn/fsn mutation produces mice with skin resembling human psoriasis. This mouse, or another mutant mouse with a similar phenotype is a further *in vivo* model to test the efficacy of topically applied therapies for psoriasis.

Another aspect of the present invention contemplates a pharmaceutical composition for topical administration which comprises a nucleic acid molecule capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation such as psoriasis and one or more pharmaceutically acceptable carriers and/or diluents. Preferably, the nucleic acid molecule is an antisense molecule to IGF-I, the IGF-I receptor or an IGFBP such as IGFBP-2 and/or IGFBP-3 or comprises a ribozyme to one or more of these targets or is a molecule suitable for co-suppression of one or more of these targets. The composition may comprise a single species of a nucleic acid molecule capable of targeting one of IGF-I, its receptor or an IGFBP, such as IGFBP-2 or IGFBP-3 or may be a multi-valent molecule capable of targeting two or more of IGF-I, its receptor or an IGFBP, such as IGFBP-2 and/or IGFBP-3.

The nucleic acid molecules may be administered in dispersions prepared in creams, ointments, oil or other suitable carrier and/or diluent such as glycerol, liquid polyethylene glycols and/or mixtures thereof. Under ordinary conditions of storage and use, these preparations may contain a preservative to prevent the growth of microorganisms.

The pharmaceutical forms suitable for topical use include sterile aqueous solutions (where water soluble) or dispersions and powders for the extemporaneous preparation of topical solutions or dispersions. In all cases, the form is preferably sterile although this is not an absolute requirement and is stable under the conditions of manufacture and storage. The carrier can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol, and the like), suitable mixtures thereof and vegetable oils. The proper fluidity can be maintained, for example, by the use of a coating such as lecithin, by the maintenance of the required particle size in the case of dispersion and by the use of superfactants. The prevention of the action of microorganism can be brought about by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal and the like. In many cases, it will be preferable to include isotonic agents, for example, sugars or sodium chloride.

Topical solutions are prepared by incorporating the nucleic acid molecule compound in the required amount in the appropriate solvent with various of the other ingredients enumerated above, as required, followed by where necessary filter sterilization.

The active agent may alternatively be administered by intravenous, subcutaneous, nasal drip, suppository, implant means amongst other suitable routes of administration including intraperitoneal, intramuscular, absorption through epithelial or mucocutaneous linings for example via nasal, oral, vaginal, rectal or gastrointestinal administration. Reference may conveniently be made to reference 24.

As used herein "pharmaceutically acceptable carriers and/or diluents" include any and all solvents, dispersion media, aqueous solutions, coatings, antibacterial and antifungal agents, isotonic and absorption delaying agents, and the like. The use of such media and agents for pharmaceutical active substances is well known in the art. Except insofar as any conventional media or agent is incompatible with the active ingredient, use thereof in the pharmaceutical compositions is contemplated. Supplementary active ingredients can also be incorporated into the compositions. Conveniently, the nucleic acid molecules of the present invention are stored in freeze-dried form and are reconstituted prior to use.

Yet another aspect of the present invention contemplates the use of a nucleic acid molecule in the manufacture of a medicament for the treatment of proliferative and/or inflammatory skin disorders or other medical disorders mediated by a growth factor. The proliferative and/or inflammatory skin disorder is generally psoriasis or other medical disorders as described above and the nucleic acid molecule targets IGF-I, the IGF-I receptor and/or an IGFBP such as IGFBP-2 and/or IGFBP-3.

Still a further aspect of the present invention contemplates an agent comprising a nucleic acid molecule as hereinbefore defined useful in the treatment of proliferative and/or inflammatory skin disorders, such as psoriasis or other medical disorder..

The present invention further contemplates the use of the genetic molecules and in particular the antisense molecules to inhibit the anti-apoptotic activity of IGF-I receptor. Such a use is appropriate for the treatment of certain cancers and as adjunct therapy for epidermal hyperplasia such as in combination with UV treatment.

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The present invention is further described by the following non-limiting Examples.

EXAMPLE 1

The differentiated human keratinocyte cell line, HaCaT [9] was used in the *in vitro* assay. Cells at passage numbers 33 to 36 were maintained as monolayer cultures in 5% v/v CO_2 at 37°C in Keratinocyte-SFM (Gibco) containing EGF and bovine pituitary extract as supplied. Media containing foetal calf serum were avoided because of the high content of IGF-I binding proteins in serum.

Feeder layer plates of lethally irradiated 3T3 fibroblasts were prepared exactly as described by Rheinwald and Green [10].

EXAMPLE 2

Cells were grown to 4 days post confluence in $2cm^2$ wells with daily medium changes of Keratinocyte-SFM, then the medium was changed to DMEM (Cytosystems, Australia), with the following additions: 25mM Hepes, 0.19% w/v, sodium bicarbonate, 0.03% w/v glutamine (Sigma Chemical Co, USA), 50IU/ml penicillin and 50µg/ml streptomycin (Flow Laboratories). After 24 hours, IGF-I or tIGF-I was added to triplicate wells, at the concentrations indicated, in 0.5ml fresh DMEM containing 0.02% v/v bovine serum albumin (Sigma molecular biology grade) and incubated for a further 21 hours. [^{3}H]-Thymidine (0.1µCi/well) was then added and the cells incubated for a further 3 hours. The medium was then aspirated and the cells washed once with ice-cold PBS and twice with ice-cold 10% v/v TCA. The TCA-precipitated monolayers were then solubilized with 0.25M NaOH (200µl/well), transferred to scintillation vials and radioactivity determined by liquid scintillation counting (Pharmacia Wallac 1410 liquid scintillation counter).

EXAMPLE 3

HaCaT conditioned medium (250µl) was concentrated by adding 750µl cold ethanol, incubating at -20°C for 2 hours and centrifuging at 16,000g for 20 min at 4°C. The resulting pellet was air dried, resuspended thoroughly in non-reducing Laemmli sample buffer, heated to 90°C for 5 minutes and separated on 12% w/v SDS-PAGE according to the method of Laemmli (1970). Separated proteins were electrophoretically transferred to nitrocellulose membrane (0.45mm, Schleicher and Schuell, Dassel, Germany) in a buffer containing 25mM Tris, 192mM glycine

and 20% v/v methanol. IGFBPs were then visualised by the procedure of Hossenlopp *et al* [11], using [^{125}I]-IGF-I, followed by autoradiography. Autoradiographs were scanned in a BioRad Model GS-670 Imaging Densitometer and band densities were determined using the Molecular Analyst program.

EXAMPLE 4

Phosphorothioate oligodeoxynucleotides were synthesised by Bresatec, Adelaide, South Australia, Australia. The following antisense sequences were used: BP3AS2, 5′- GCG CCC GCT GCA TGA CGC CTG CAA C -3′ (<400>4), a 25mer complementary to the start codon region of the human IGFBP-3 mRNA; BP3AS3, 5′- CGG GCG GCT CAC CTG GAG CTG GCG -3′ (<400>5), a 24mer complementary to the exon 1/intron 1 splice site; BP3AS4, 5′- AGG CGG CTG ACG GCA CTA -3′(<400>6), an 18mer complementary to a region of the coding sequence lacking RNA secondary structure and oligonucleotide-dimer formation (using the computer software "OLIGO for PC"). Since BP3AS4 was found to be ineffective at inhibiting IGFBP-3 synthesis, it was used as a control. The following additional control oligonucleotide sequences were used: BP3S, 5′- CAG GCG TCA TGC AGC GGG C -3′ (<400>7), an 18mer sense control sequence equivalent to the start codon region; BP3AS2NS, 5′- CGG AGA TGC CGC ATG CCA GCG CAG G -3′ (<400>8), a 25mer randomised sequence with the same GC content as BP3AS2; BP3AS4NS, 5′- GAC AGC GTC GGA GCG ATC -3′ (<400>9), an 18mer randomised sequence with the same GC content as BP3AS4NS. Design of the oligonucleotides was based on the human IGFBP-3 cDNA sequence of Spratt *et al* [12].

Cells were grown to one day post confluence in 2cm^2 wells with daily medium changes of 0.5ml Keratinocyte-SFM, then subjected to daily medium changes of Keratinocyte-SFM for a further 4 days. Daily additions of 0.5ml fresh Keratinocyte-SFM were then continued for a further 7 days, except that at the time of medium addition, 5µl oligonucleotide in PBS was added to give the final concentrations indicated, then the wells were shaken to mix the oligonucleotide. After the final addition, cells were incubated for 24 hours and the medium collected for assay of IGFBPs. Cells were then counted after trypsinisation in a Coulter Industrial D Counter, Coulter Bedfordshire, UK. Cell numbers after oligonucleotide treatment differed by less than 10%.

EXAMPLE 5

HaCaT cells secrete mainly IGFBP-3 (>95%), with the only other IGFBP detectable in HaCaT conditioned medium being IGFBP-4 (<5%). The effect on IGFBP-3 and IGFBP-4 synthesis of antisense oligonucleotides at two concentrations, 5μM and 0.5μM, was tested. Two oligonucleotides were used, BP3AS2 and BP3AS3, directed against the start site and the intron 1/exon 1 splice site, respectively of the IGFBP-3 mRNA. As a control, a sense oligonucleotide corresponding to the start site was used. As shown in Figures 4A and 4B, all oligonucleotides at 5μM caused a significant reduction of IGFBP-3 synthesis compared with untreated cells, however, the two antisense oligonucleotides inhibited IGFBP-3 synthesis of approximately 50% compared to the sense control (Figure 4B). The antisense oligonucleotide directed to the start codon appeared to be more effective of the two, the difference being more apparent at the lower concentration of 0.5μM. The cells of IGFBP-4 secreted by the HaCaT cells make photographic reproduction of the bands on Western ligand blots difficult, however densitometry measurements provide adequate relative quantitation. This resulted in the significant observation that IGFBP-4 levels were unaffected by oligonucleotide addition to the cells, suggesting that the observed inhibitory effects on IGFBP-3 are specific.

To further investigate the inhibitory effects of the more effective of the two antisense oligonucleotides, BP3AS2, inhibition by this oligonucleotide at 0.5μM was compared with a number of control oligonucleotides, including one antisense oligonucleotide to IGFBP-3 that had proved to be ineffective at 0.5μM. As shown in Figures 5A and 5B, BP3AS2 was again inhibitory, resulting in levels of IGFBP-3 of approximately 50% of the most non-specifically inhibitory control oligonucleotide, the randomised equivalent of BP3AS2. The other control oligonucleotides caused no reduction in IGFBP-3 levels at 0.5μM, compared to untreated cells. Of possible significance is the fact that this control oligonucleotide, BP3AS2NS, like BP3AS2 itself, has the highest potential T_m of the three control oligonucleotides used in this experiment, enhancing the probability of non-specific base pairing with non-target mRNAs. However, the lack of inhibition of IGFBP-4 secretion by BP3AS2 suggests that this oligonucleotide is selective even compared with the most closely related protein likely to be present in this cell line.

EXAMPLE 6

Antisense oligonucleotides to IGFBP2 may be selected from molecules capable of interacting with one or more of the following sense oligonucleotides:

ATTCGGGGCGAGGGA	CGCAGGGCCGTGCAC	CCGCGCCGCGCTGCC
TTCGGGGCGAGGGAG	GCAGGGCCGTGCACC	CGCGCCGCGCTGCCG
TCGGGGCGAGGGAGG	CAGGGCCGTGCACCT	GCGCCGCGCTGCCGA
CGGGGCGAGGGAGGA	AGGGCCGTGCACCTG	CGCCGCGCTGCCGAC
GGGGCGAGGGAGGAG	GGGCCGTGCACCTGC	GCCGCGCTGCCGACC
GGGCGAGGGAGGAGG	GGCCGTGCACCTGCC	CCGCGCTGCCGACCG
GGCGAGGGAGGAGGA	GCCGTGCACCTGCCC	CGCGCTGCCGACCGC
GCGAGGGAGGAGGAA	CCGTGCACCTGCCCG	GCGCTGCCGACCGCC
CGAGGGAGGAGGAAG	CGTGCACCTGCCCGC	CGCTGCCGACCGCCA
GAGGGAGGAGGAAGA	GTGCACCTGCCCGCC	GCTGCCGACCGCCAG
AGGGAGGAGGAAGAA	TGCACCTGCCCGCCC	CTGCCGACCGCCAGC
GGGAGGAGGAAGAAG	GCACCTGCCCGCCCG	TGCCGACCGCCAGCA
GGAGGAGGAAGAAGC	CACCTGCCCGCCCGC	GCCGACCGCCAGCAT
GAGGAGGAAGAAGCG	ACCTGCCCGCCCGCC	CCGACCGCCAGCATG
AGGAGGAAGAAGCGG	CCTGCCCGCCCGCCC	CGACCGCCAGCATGC
GGAGGAAGAAGCGGA	CTGCCCGCCCGCCCG	GACCGCCAGCATGCT
GAGGAAGAAGCGGAG	TGCCCGCCCGCCCGC	ACCGCCAGCATGCTG
AGGAAGAAGCGGAGG	GCCCGCCCGCCCGCT	CCGCCAGCATGCTGC
GGAAGAAGCGGAGGA	CCCGCCCGCCCGCTC	CGCCAGCATGCTGCC
GAAGAAGCGGAGGAG	CCGCCCGCCCGCTCG	GCCAGCATGCTGCCG
AAGAAGCGGAGGAGG	CGCCCGCCCGCTCGC	CCAGCATGCTGCCGA
AGAAGCGGAGGAGGC	GCCCGCCCGCTCGCT	CAGCATGCTGCCGAG
GAAGCGGAGGAGGCG	CCCGCCCGCTCGCTC	AGCATGCTGCCGAGA
AAGCGGAGGAGGCGG	CCGCCCGCTCGCTCG	GCATGCTGCCGAGAG
AGCGGAGGAGGCGGC	CGCCCGCTCGCTCGC	CATGCTGCCGAGAGT
GCGGAGGAGGCGGCT	GCCCGCTCGCTCGCT	ATGCTGCCGAGAGTG
CGGAGGAGGCGGCTC	CCCGCTCGCTCGCTC	TGCTGCCGAGAGTGG
GGAGGAGGCGGCTCC	CCGCTCGCTCGCTCG	GCTGCCGAGAGTGGG
GAGGAGGCGGCTCCC	CGCTCGCTCGCTCGC	CTGCCGAGAGTGGGC
AGGAGGCGGCTCCCG	GCTCGCTCGCTCGCC	TGCCGAGAGTGGGCT
GGAGGCGGCTCCCGC	CTCGCTCGCTCGCCC	GCCGAGAGTGGGCTG
GAGGCGGCTCCCGCT	TCGCTCGCTCGCCCG	CCGAGAGTGGGCTGC
AGGCGGCTCCCGCTC	CGCTCGCTCGCCCGC	CGAGAGTGGGCTGCC
GGCGGCTCCCGCTCG	GCTCGCTCGCCCGCC	GAGAGTGGGCTGCCC
GCGGCTCCCGCTCGC	CTCGCTCGCCCGCCG	AGAGTGGGCTGCCCC
CGGCTCCCGCTCGCA	TCGCTCGCCCGCCGC	GAGTGGGCTGCCCCG
GGCTCCCGCTCGCAG	CGCTCGCCCGCCGCG	AGTGGGCTGCCCCGC
GCTCCCGCTCGCAGG	GCTCGCCCGCCGCGC	GTGGGCTGCCCCGCG
CTCCCGCTCGCAGGG	CTCGCCCGCCGCGCC	TGGGCTGCCCCGCGC
TCCCGCTCGCAGGGC	TCGCCCGCCGCGCCG	GGGCTGCCCCGCGCT
CCCGCTCGCAGGGCC	CGCCCGCCGCGCCGC	GGCTGCCCCGCGCTG
CCGCTCGCAGGGCCG	GCCCGCCGCGCCGCG	GCTGCCCCGCGCTGC
CGCTCGCAGGGCCGT	CCCGCCGCGCCGCGC	CTGCCCCGCGCTGCC
GCTCGCAGGGCCGTG	CCGCCGCGCCGCGCT	TGCCCCGCGCTGCCG
CTCGCAGGGCCGTGC	CGCCGCGCCGCGCTG	GCCCCGCGCTGCCGC
TCGCAGGGCCGTGCA	GCCGCGCCGCGCTGC	CCCCGCGCTGCCGCT

	CCCGCGCTGCCGCTG	CTGCTGCTACTGGGC	CTGTTCCGCTGCCCG
	CCGCGCTGCCGCTGC	TGCTGCTACTGGGCG	TGTTCCGCTGCCCGC
	CGCGCTGCCGCTGCC	GCTGCTACTGGGCGC	GTTCCGCTGCCCGCC
	GCGCTGCCGCTGCCG	CTGCTACTGGGCGCG	TTCCGCTGCCCGCCC
5	CGCTGCCGCTGCCGC	TGCTACTGGGCGCGA	TCCGCTGCCCGCCCT
	GCTGCCGCTGCCGCC	GCTACTGGGCGCGAG	CCGCTGCCCGCCCTG
	CTGCCGCTGCCGCCG	CTACTGGGCGCGAGT	CGCTGCCCGCCCTGC
	TGCCGCTGCCGCCGC	TACTGGGCGCGAGTG	GCTGCCCGCCCTGCA
	GCCGCTGCCGCCGCC	ACTGGGCGCGAGTGG	CTGCCCGCCCTGCAC
10	CCGCTGCCGCCGCCG	CTGGGCGCGAGTGGC	TGCCCGCCCTGCACA
	CGCTGCCGCCGCCGC	TGGGCGCGAGTGGCG	GCCCGCCCTGCACAC
	GCTGCCGCCGCCGCC	GGGCGCGAGTGGCGG	CCCGCCCTGCACACC
	CTGCCGCCGCCGCCG	GGCGCGAGTGGCGGC	CCGCCCTGCACACCC
	TGCCGCCGCCGCCGC	GCGCGAGTGGCGGCG	CGCCCTGCACACCCG
15	GCCGCCGCCGCCGCT	CGCGAGTGGCGGCGG	GCCCTGCACACCCGA
	CCGCCGCCGCCGCTG	GCGAGTGGCGGCGGC	CCCTGCACACCCGAG
	CGCCGCCGCCGCTGC	CGAGTGGCGGCGGCG	CCTGCACACCCGAGC
	GCCGCCGCCGCTGCT	GAGTGGCGGCGGCGG	CTGCACACCCGAGCG
	CCGCCGCCGCTGCTG	AGTGGCGGCGGCGGC	TGCACACCCGAGCGC
20	CGCCGCCGCTGCTGC	GTGGCGGCGGCGGCG	GCACACCCGAGCGCC
	GCCGCCGCTGCTGCC	TGGCGGCGGCGGCGG	CACACCCGAGCGCCT
	CCGCCGCTGCTGCCG	GGCGGCGGCGGCGGG	ACACCCGAGCGCCTG
	CGCCGCTGCTGCCGC	GCGGCGGCGGCGGGG	CACCCGAGCGCCTGG
	GCCGCTGCTGCCGCT	CGGCGGCGGCGGGGC	ACCCGAGCGCCTGGC
25	CCGCTGCTGCCGCTG	GGCGGCGGCGGGGCG	CCCGAGCGCCTGGCC
	CGCTGCTGCCGCTGC	GCGGCGGCGGGGCGC	CCGAGCGCCTGGCCG
	GCTGCTGCCGCTGCT	CGGCGGCGGGGCGCG	CGAGCGCCTGGCCGC
	CTGCTGCCGCTGCTG	GGCGGCGGGGCGCGC	GAGCGCCTGGCCGCC
	TGCTGCCGCTGCTGC	GCGGCGGGGCGCGCG	AGCGCCTGGCCGCCT
30	GCTGCCGCTGCTGCC	CGGCGGGGCGCGCGC	GCGCCTGGCCGCCTG
	CTGCCGCTGCTGCCG	GGCGGGGCGCGCGCG	CGCCTGGCCGCCTGC
	TGCCGCTGCTGCCGC	GCGGGGCGCGCGCGG	GCCTGGCCGCCTGCG
	GCCGCTGCTGCCGCT	CGGGGCGCGCGCGGA	CCTGGCCGCCTGCGG
	CCGCTGCTGCCGCTG	GGGGCGCGCGCGGAG	CTGGCCGCCTGCGGG
35	CGCTGCTGCCGCTGC	GGGCGCGCGCGGAGG	TGGCCGCCTGCGGGC
	GCTGCTGCCGCTGCT	GGCGCGCGCGGAGGT	GGCCGCCTGCGGGCC
	CTGCTGCCGCTGCTG	GCGCGCGCGGAGGTG	GCCGCCTGCGGGCCC
	TGCTGCCGCTGCTGC	CGCGCGCGGAGGTGC	CCGCCTGCGGGCCCC
	GCTGCCGCTGCTGCT	GCGCGCGGAGGTGCT	CGCCTGCGGGCCCCC
40	CTGCCGCTGCTGCTG	CGCGCGGAGGTGCTG	GCCTGCGGGCCCCCG
	TGCCGCTGCTGCTGC	GCGCGGAGGTGCTGT	CCTGCGGGCCCCCGC
	GCCGCTGCTGCTGCT	CGCGGAGGTGCTGTT	CTGCGGGCCCCCGCC
	CCGCTGCTGCTGCTG	GCGGAGGTGCTGTTC	TGCGGGCCCCCGCCG
	CGCTGCTGCTGCTGC	CGGAGGTGCTGTTCC	GCGGGCCCCCGCCGG
45	GCTGCTGCTGCTGCT	GGAGGTGCTGTTCCG	CGGGCCCCCGCCGGT
	CTGCTGCTGCTGCTA	GAGGTGCTGTTCCGC	GGGCCCCCGCCGGTT
	TGCTGCTGCTGCTAC	AGGTGCTGTTCCGCT	GGCCCCCGCCGGTTG
	GCTGCTGCTGCTACT	GGTGCTGTTCCGCTG	GCCCCCGCCGGTTGC
	CTGCTGCTGCTACTG	GTGCTGTTCCGCTGC	CCCCCGCCGGTTGCG
50	TGCTGCTGCTACTGG	TGCTGTTCCGCTGCC	CCCCGCCGGTTGCGC
	GCTGCTGCTACTGGG	GCTGTTCCGCTGCCC	CCCGCCGGTTGCGCC

	CCGCCGGTTGCGCCG	ATGCCATGCGCGGAG	TGCGCCCGGCTGGAG
	CGCCGGTTGCGCCGC	TGCCATGCGCGGAGC	GCGCCCGGCTGGAGG
	GCCGGTTGCGCCGCC	GCCATGCGCGGAGCT	CGCCCGGCTGGAGGG
	CCGGTTGCGCCGCCC	CCATGCGCGGAGCTC	GCCCGGCTGGAGGGC
5	CGGTTGCGCCGCCCG	CATGCGCGGAGCTCG	CCCGGCTGGAGGGCG
	GGTTGCGCCGCCCGC	ATGCGCGGAGCTCGT	CCGGCTGGAGGGCGA
	GTTGCGCCGCCCGCC	TGCGCGGAGCTCGTC	CGGCTGGAGGGCGAG
	TTGCGCCGCCCGCCG	GCGCGGAGCTCGTCC	GGCTGGAGGGCGAGG
	TGCGCCGCCCGCCGC	CGCGGAGCTCGTCCG	GCTGGAGGGCGAGGC
10	GCGCCGCCCGCCGCG	GCGGAGCTCGTCCGG	CTGGAGGGCGAGGCG
	CGCCGCCCGCCGCGG	CGGAGCTCGTCCGGG	TGGAGGGCGAGGCGT
	GCCGCCCGCCGCGGT	GGAGCTCGTCCGGGA	GGAGGGCGAGGCGTG
	CCGCCCGCCGCGGTG	GAGCTCGTCCGGGAG	GAGGGCGAGGCGTGC
	CGCCCGCCGCGGTGG	AGCTCGTCCGGGAGC	AGGGCGAGGCGTGCG
15	GCCCGCCGCGGTGGC	GCTCGTCCGGGAGCC	GGGCGAGGCGTGCGG
	CCCGCCGCGGTGGCC	CTCGTCCGGGAGCCG	GGCGAGGCGTGCGGC
	CCGCCGCGGTGGCCG	TCGTCCGGGAGCCGG	GCGAGGCGTGCGGCG
	CGCCGCGGTGGCCGC	CGTCCGGGAGCCGGG	CGAGGCGTGCGGCGT
	GCCGCGGTGGCCGCA	GTCCGGGAGCCGGGC	GAGGCGTGCGGCGTC
20	CCGCGGTGGCCGCAG	TCCGGGAGCCGGGCT	AGGCGTGCGGCGTCT
	CGCGGTGGCCGCAGT	CCGGGAGCCGGGCTG	GGCGTGCGGCGTCTA
	GCGGTGGCCGCAGTG	CGGGAGCCGGGCTGC	GCGTGCGGCGTCTAC
	CGGTGGCCGCAGTGG	GGGAGCCGGGCTGCG	CGTGCGGCGTCTACA
	GGTGGCCGCAGTGGC	GGAGCCGGGCTGCGG	GTGCGGCGTCTACAC
25	GTGGCCGCAGTGGCC	GAGCCGGGCTGCGGC	TGCGGCGTCTACACC
	TGGCCGCAGTGGCCG	AGCCGGGCTGCGGCT	GCGGCGTCTACACCC
	GGCCGCAGTGGCCGG	GCCGGGCTGCGGCTG	CGGCGTCTACACCCC
	GCCGCAGTGGCCGGA	CCGGGCTGCGGCTGC	GGCGTCTACACCCCG
	CCGCAGTGGCCGGAG	CGGGCTGCGGCTGCT	GCGTCTACACCCCGC
30	CGCAGTGGCCGGAGG	GGGCTGCGGCTGCTG	CGTCTACACCCCGCG
	GCAGTGGCCGGAGGC	GGCTGCGGCTGCTGC	GTCTACACCCCGCGC
	CAGTGGCCGGAGGCG	GCTGCGGCTGCTGCT	TCTACACCCCGCGCT
	AGTGGCCGGAGGCGC	CTGCGGCTGCTGCTC	CTACACCCCGCGCTG
	GTGGCCGGAGGCGCC	TGCGGCTGCTGCTCG	TACACCCCGCGCTGC
35	TGGCCGGAGGCGCCC	GCGGCTGCTGCTCGG	ACACCCCGCGCTGCG
	GGCCGGAGGCGCCCG	CGGCTGCTGCTCGGT	CACCCCGCGCTGCGG
	GCCGGAGGCGCCCGC	GGCTGCTGCTCGGTG	ACCCCGCGCTGCGGC
	CCGGAGGCGCCCGCA	GCTGCTGCTCGGTGT	CCCCGCGCTGCGGCC
	CGGAGGCGCCCGCAT	CTGCTGCTCGGTGTG	CCCGCGCTGCGGCCA
40	GGAGGCGCCCGCATG	TGCTGCTCGGTGTGC	CCGCGCTGCGGCCAG
	GAGGCGCCCGCATGC	GCTGCTCGGTGTGCG	CGCGCTGCGGCCAGG
	AGGCGCCCGCATGCC	CTGCTCGGTGTGCGC	GCGCTGCGGCCAGGG
	GGCGCCCGCATGCCA	TGCTCGGTGTGCGCC	CGCTGCGGCCAGGGG
	GCGCCCGCATGCCAT	GCTCGGTGTGCGCCC	GCTGCGGCCAGGGGC
45	CGCCCGCATGCCATG	CTCGGTGTGCGCCCG	CTGCGGCCAGGGGCT
	GCCCGCATGCCATGC	TCGGTGTGCGCCCGG	TGCGGCCAGGGGCTG
	CCCGCATGCCATGCG	CGGTGTGCGCCCGGC	GCGGCCAGGGGCTGC
	CCGCATGCCATGCGC	GGTGTGCGCCCGGCT	CGGCCAGGGGCTGCG
	CGCATGCCATGCGCG	GTGTGCGCCCGGCTG	GGCCAGGGGCTGCGC
50	GCATGCCATGCGCGG	TGTGCGCCCGGCTGG	GCCAGGGGCTGCGCT
	CATGCCATGCGCGGA	GTGCGCCCGGCTGGA	CCAGGGGCTGCGCTG

CAGGGGCTGCGCTGC	CTGGTCATGGGCGAG	GCCAGCCCGGAGCAG
AGGGGCTGCGCTGCT	TGGTCATGGGCGAGG	CCAGCCCGGAGCAGG
GGGGCTGCGCTGCTA	GGTCATGGGCGAGGG	CAGCCCGGAGCAGGT
GGGCTGCGCTGCTAT	GTCATGGGCGAGGGC	AGCCCGGAGCAGGTT
GGCTGCGCTGCTATC	TCATGGGCGAGGGCA	GCCCGGAGCAGGTTG
GCTGCGCTGCTATCC	CATGGGCGAGGGCAC	CCCGGAGCAGGTTGC
CTGCGCTGCTATCCC	ATGGGCGAGGGCACT	CCGGAGCAGGTTGCA
TGCGCTGCTATCCCC	TGGGCGAGGGCACTT	CGGAGCAGGTTGCAG
GCGCTGCTATCCCCA	GGGCGAGGGCACTTG	GGAGCAGGTTGCAGA
CGCTGCTATCCCCAC	GGCGAGGGCACTTGT	GAGCAGGTTGCAGAC
GCTGCTATCCCCACC	GCGAGGGCACTTGTG	AGCAGGTTGCAGACA
CTGCTATCCCCACCC	CGAGGGCACTTGTGA	GCAGGTTGCAGACAA
TGCTATCCCCACCCG	GAGGGCACTTGTGAG	CAGGTTGCAGACAAT
GCTATCCCCACCCGG	AGGGCACTTGTGAGA	AGGTTGCAGACAATG
CTATCCCCACCCGGG	GGGCACTTGTGAGAA	GGTTGCAGACAATGG
TATCCCCACCCGGGC	GGCACTTGTGAGAAG	GTTGCAGACAATGGC
ATCCCCACCCGGGCT	GCACTTGTGAGAAGC	TTGCAGACAATGGCG
TCCCCACCCGGGCTC	CACTTGTGAGAAGCG	TGCAGACAATGGCGA
CCCCACCCGGGCTCC	ACTTGTGAGAAGCGC	GCAGACAATGGCGAT
CCCACCCGGGCTCCG	CTTGTGAGAAGCGCC	CAGACAATGGCGATG
CCACCCGGGCTCCGA	TTGTGAGAAGCGCCG	AGACAATGGCGATGA
CACCCGGGCTCCGAG	TGTGAGAAGCGCCGG	GACAATGGCGATGAC
ACCCGGGCTCCGAGC	GTGAGAAGCGCCGGG	ACAATGGCGATGACC
CCCGGGCTCCGAGCT	TGAGAAGCGCCGGGA	CAATGGCGATGACCA
CCGGGCTCCGAGCTG	GAGAAGCGCCGGGAC	AATGGCGATGACCAC
CGGGCTCCGAGCTGC	AGAAGCGCCGGGACG	ATGGCGATGACCACT
GGGCTCCGAGCTGCC	GAAGCGCCGGGACGC	TGGCGATGACCACTC
GGCTCCGAGCTGCCC	AAGCGCCGGGACGCC	GGCGATGACCACTCA
GCTCCGAGCTGCCCC	AGCGCCGGGACGCCG	GCGATGACCACTCAG
CTCCGAGCTGCCCCT	GCGCCGGGACGCCGA	CGATGACCACTCAGA
TCCGAGCTGCCCCTG	CGCCGGGACGCCGAG	GATGACCACTCAGAA
CCGAGCTGCCCCTGC	GCCGGGACGCCGAGT	ATGACCACTCAGAAG
CGAGCTGCCCCTGCA	CCGGGACGCCGAGTA	TGACCACTCAGAAGG
GAGCTGCCCCTGCAG	CGGGACGCCGAGTAT	GACCACTCAGAAGGA
AGCTGCCCCTGCAGG	GGGACGCCGAGTATG	ACCACTCAGAAGGAG
GCTGCCCCTGCAGGC	GGACGCCGAGTATGG	CCACTCAGAAGGAGG
CTGCCCCTGCAGGCG	GACGCCGAGTATGGC	CACTCAGAAGGAGGC
TGCCCCTGCAGGCGC	ACGCCGAGTATGGCG	ACTCAGAAGGAGGCC
GCCCCTGCAGGCGCT	CGCCGAGTATGGCGC	CTCAGAAGGAGGCCT
CCCCTGCAGGCGCTG	GCCGAGTATGGCGCC	TCAGAAGGAGGCCTG
CCCTGCAGGCGCTGG	CCGAGTATGGCGCCA	CAGAAGGAGGCCTGG
CCTGCAGGCGCTGGT	CGAGTATGGCGCCAG	AGAAGGAGGCCTGGT
CTGCAGGCGCTGGTC	GAGTATGGCGCCAGC	GAAGGAGGCCTGGTG
TGCAGGCGCTGGTCA	AGTATGGCGCCAGCC	AAGGAGGCCTGGTGG
GCAGGCGCTGGTCAT	GTATGGCGCCAGCCC	AGGAGGCCTGGTGGA
CAGGCGCTGGTCATG	TATGGCGCCAGCCCG	GGAGGCCTGGTGGAG
AGGCGCTGGTCATGG	ATGGCGCCAGCCCGG	GAGGCCTGGTGGAGA
GGCGCTGGTCATGGG	TGGCGCCAGCCCGGA	AGGCCTGGTGGAGAA
GCGCTGGTCATGGGC	GGCGCCAGCCCGGAG	GGCCTGGTGGAGAAC
CGCTGGTCATGGGCG	GCGCCAGCCCGGAGC	GCCTGGTGGAGAACC
GCTGGTCATGGGCGA	CGCCAGCCCGGAGCA	CCTGGTGGAGAACCA

CTGGTGGAGAACCAC	AGTGCTGGCCGGAAG	CGGGAGAAGGTCACT
TGGTGGAGAACCACG	GTGCTGGCCGGAAGC	GGGAGAAGGTCACTG
GGTGGAGAACCACGT	TGCTGGCCGGAAGCC	GGAGAAGGTCACTGA
GTGGAGAACCACGTG	GCTGGCCGGAAGCCC	GAGAAGGTCACTGAG
TGGAGAACCACGTGG	CTGGCCGGAAGCCCC	AGAAGGTCACTGAGC
GGAGAACCACGTGGA	TGGCCGGAAGCCCCT	GAAGGTCACTGAGCA
GAGAACCACGTGGAC	GGCCGGAAGCCCCTC	AAGGTCACTGAGCAG
AGAACCACGTGGACA	GCCGGAAGCCCCTCA	AGGTCACTGAGCAGC
GAACCACGTGGACAG	CCGGAAGCCCCTCAA	GGTCACTGAGCAGCA
AACCACGTGGACAGC	CGGAAGCCCCTCAAG	GTCACTGAGCAGCAC
ACCACGTGGACAGCA	GGAAGCCCCTCAAGT	TCACTGAGCAGCACC
CCACGTGGACAGCAC	GAAGCCCCTCAAGTC	CACTGAGCAGCACCG
CACGTGGACAGCACC	AAGCCCCTCAAGTCG	ACTGAGCAGCACCGG
ACGTGGACAGCACCA	AGCCCCTCAAGTCGG	CTGAGCAGCACCGGC
CGTGGACAGCACCAT	GCCCCTCAAGTCGGG	TGAGCAGCACCGGCA
GTGGACAGCACCATG	CCCCTCAAGTCGGGT	GAGCAGCACCGGCAG
TGGACAGCACCATGA	CCCTCAAGTCGGGTA	AGCAGCACCGGCAGA
GGACAGCACCATGAA	CCTCAAGTCGGGTAT	GCAGCACCGGCAGAT
GACAGCACCATGAAC	CTCAAGTCGGGTATG	CAGCACCGGCAGATG
ACAGCACCATGAACA	TCAAGTCGGGTATGA	AGCACCGGCAGATGG
CAGCACCATGAACAT	CAAGTCGGGTATGAA	GCACCGGCAGATGGG
AGCACCATGAACATG	AAGTCGGGTATGAAG	CACCGGCAGATGGGC
GCACCATGAACATGT	AGTCGGGTATGAAGG	ACCGGCAGATGGGCA
CACCATGAACATGTT	GTCGGGTATGAAGGA	CCGGCAGATGGGCAA
ACCATGAACATGTTG	TCGGGTATGAAGGAG	CGGCAGATGGGCAAG
CCATGAACATGTTGG	CGGGTATGAAGGAGC	GGCAGATGGGCAAGG
CATGAACATGTTGGG	GGGTATGAAGGAGCT	GCAGATGGGCAAGGG
ATGAACATGTTGGGC	GGTATGAAGGAGCTG	CAGATGGGCAAGGGT
TGAACATGTTGGGCG	GTATGAAGGAGCTGG	AGATGGGCAAGGGTG
GAACATGTTGGGCGG	TATGAAGGAGCTGGC	GATGGGCAAGGGTGG
AACATGTTGGGCGGG	ATGAAGGAGCTGGCC	ATGGGCAAGGGTGGC
ACATGTTGGGCGGGG	TGAAGGAGCTGGCCG	TGGGCAAGGGTGGCA
CATGTTGGGCGGGGG	GAAGGAGCTGGCCGT	GGGCAAGGGTGGCAA
ATGTTGGGCGGGGGA	AAGGAGCTGGCCGTG	GGCAAGGGTGGCAAG
TGTTGGGCGGGGGAG	AGGAGCTGGCCGTGT	GCAAGGGTGGCAAGC
GTTGGGCGGGGGAGG	GGAGCTGGCCGTGTT	CAAGGGTGGCAAGCA
TTGGGCGGGGGAGGC	GAGCTGGCCGTGTTC	AAGGGTGGCAAGCAT
TGGGCGGGGGAGGCA	AGCTGGCCGTGTTCC	AGGGTGGCAAGCATC
GGGCGGGGGAGGCAG	GCTGGCCGTGTTCCG	GGGTGGCAAGCATCA
GGCGGGGGAGGCAGT	CTGGCCGTGTTCCGG	GGTGGCAAGCATCAC
GCGGGGGAGGCAGTG	TGGCCGTGTTCCGGG	GTGGCAAGCATCACC
CGGGGGAGGCAGTGC	GGCCGTGTTCCGGGA	TGGCAAGCATCACCT
GGGGGAGGCAGTGCT	GCCGTGTTCCGGGAG	GGCAAGCATCACCTT
GGGGAGGCAGTGCTG	CCGTGTTCCGGGAGA	GCAAGCATCACCTTG
GGGAGGCAGTGCTGG	CGTGTTCCGGGAGAA	CAAGCATCACCTTGG
GGAGGCAGTGCTGGC	GTGTTCCGGGAGAAG	AAGCATCACCTTGGC
GAGGCAGTGCTGGCC	TGTTCCGGGAGAAGG	AGCATCACCTTGGCC
AGGCAGTGCTGGCCG	GTTCCGGGAGAAGGT	GCATCACCTTGGCCT
GGCAGTGCTGGCCGG	TTCCGGGAGAAGGTC	CATCACCTTGGCCTG
GCAGTGCTGGCCGGA	TCCGGGAGAAGGTCA	ATCACCTTGGCCTGG
CAGTGCTGGCCGGAA	CCGGGAGAAGGTCAC	TCACCTTGGCCTGGA

CACCTTGGCCTGGAG	CCCTGCCAACAGGAA	CTTCCGGATGAGCGG
ACCTTGGCCTGGAGG	CCTGCCAACAGGAAC	TTCCGGATGAGCGGG
CCTTGGCCTGGAGGA	CTGCCAACAGGAACT	TCCGGATGAGCGGGG
CTTGGCCTGGAGGAG	TGCCAACAGGAACTG	CCGGATGAGCGGGGC
TTGGCCTGGAGGAGC	GCCAACAGGAACTGG	CGGATGAGCGGGGCC
TGGCCTGGAGGAGCC	CCAACAGGAACTGGA	GGATGAGCGGGGCCC
GGCCTGGAGGAGCCC	CAACAGGAACTGGAC	GATGAGCGGGGCCCT
GCCTGGAGGAGCCCA	AACAGGAACTGGACC	ATGAGCGGGGCCCTC
CCTGGAGGAGCCCAA	ACAGGAACTGGACCA	TGAGCGGGGCCCTCT
CTGGAGGAGCCCAAG	CAGGAACTGGACCAG	GAGCGGGGCCCTCTG
TGGAGGAGCCCAAGA	AGGAACTGGACCAGG	AGCGGGGCCCTCTGG
GGAGGAGCCCAAGAA	GGAACTGGACCAGGT	GCGGGGCCCTCTGGA
GAGGAGCCCAAGAAG	GAACTGGACCAGGTC	CGGGGCCCTCTGGAG
AGGAGCCCAAGAAGC	AACTGGACCAGGTCC	GGGGCCCTCTGGAGC
GGAGCCCAAGAAGCT	ACTGGACCAGGTCCT	GGGCCCTCTGGAGCA
GAGCCCAAGAAGCTG	CTGGACCAGGTCCTG	GGCCCTCTGGAGCAC
AGCCCAAGAAGCTGC	TGGACCAGGTCCTGG	GCCCTCTGGAGCACC
GCCCAAGAAGCTGCG	GGACCAGGTCCTGGA	CCCTCTGGAGCACCT
CCCAAGAAGCTGCGA	GACCAGGTCCTGGAG	CCTCTGGAGCACCTC
CCAAGAAGCTGCGAC	ACCAGGTCCTGGAGC	CTCTGGAGCACCTCT
CAAGAAGCTGCGACC	CCAGGTCCTGGAGCG	TCTGGAGCACCTCTA
AAGAAGCTGCGACCA	CAGGTCCTGGAGCGG	CTGGAGCACCTCTAC
AGAAGCTGCGACCAC	AGGTCCTGGAGCGGA	TGGAGCACCTCTACT
GAAGCTGCGACCACC	GGTCCTGGAGCGGAT	GGAGCACCTCTACTC
AAGCTGCGACCACCC	GTCCTGGAGCGGATC	GAGCACCTCTACTCC
AGCTGCGACCACCCC	TCCTGGAGCGGATCT	AGCACCTCTACTCCC
GCTGCGACCACCCCC	CCTGGAGCGGATCTC	GCACCTCTACTCCCT
CTGCGACCACCCCCT	CTGGAGCGGATCTCC	CACCTCTACTCCCTG
TGCGACCACCCCCTG	TGGAGCGGATCTCCA	ACCTCTACTCCCTGC
GCGACCACCCCCTGC	GGAGCGGATCTCCAC	CCTCTACTCCCTGCA
CGACCACCCCCTGCC	GAGCGGATCTCCACC	CTCTACTCCCTGCAC
GACCACCCCCTGCCA	AGCGGATCTCCACCA	TCTACTCCCTGCACA
ACCACCCCCTGCCAG	GCGGATCTCCACCAT	CTACTCCCTGCACAT
CCACCCCCTGCCAGG	CGGATCTCCACCATG	TACTCCCTGCACATC
CACCCCCTGCCAGGA	GGATCTCCACCATGC	ACTCCCTGCACATCC
ACCCCCTGCCAGGAC	GATCTCCACCATGCG	CTCCCTGCACATCCC
CCCCCTGCCAGGACT	ATCTCCACCATGCGC	TCCCTGCACATCCCC
CCCCTGCCAGGACTC	TCTCCACCATGCGCC	CCCTGCACATCCCCA
CCCTGCCAGGACTCC	CTCCACCATGCGCCT	CCTGCACATCCCCAA
CCTGCCAGGACTCCC	TCCACCATGCGCCTT	CTGCACATCCCCAAC
CTGCCAGGACTCCCT	CCACCATGCGCCTTC	TGCACATCCCCAACT
TGCCAGGACTCCCTG	CACCATGCGCCTTCC	GCACATCCCCAACTG
GCCAGGACTCCCTGC	ACCATGCGCCTTCCG	CACATCCCCAACTGT
CCAGGACTCCCTGCC	CCATGCGCCTTCCGG	ACATCCCCAACTGTG
CAGGACTCCCTGCCA	CATGCGCCTTCCGGA	CATCCCCAACTGTGA
AGGACTCCCTGCCAA	ATGCGCCTTCCGGAT	ATCCCCAACTGTGAC
GGACTCCCTGCCAAC	TGCGCCTTCCGGATG	TCCCCAACTGTGACA
GACTCCCTGCCAACA	GCGCCTTCCGGATGA	CCCCAACTGTGACAA
ACTCCCTGCCAACAG	CGCCTTCCGGATGAG	CCCAACTGTGACAAG
CTCCCTGCCAACAGG	GCCTTCCGGATGAGC	CCAACTGTGACAAGC
TCCCTGCCAACAGGA	CCTTCCGGATGAGCG	CAACTGTGACAAGCA

	AACTGTGACAAGCAT	AACGGGCAGCGTGGG	ATCCAGGGAGCCCCC
	ACTGTGACAAGCATG	ACGGGCAGCGTGGGG	TCCAGGGAGCCCCCA
	CTGTGACAAGCATGG	CGGGCAGCGTGGGGA	CCAGGGAGCCCCCAC
	TGTGACAAGCATGGC	GGGCAGCGTGGGGAG	CAGGGAGCCCCCACC
5	GTGACAAGCATGGCC	GGCAGCGTGGGGAGT	AGGGAGCCCCCACCA
	TGACAAGCATGGCCT	GCAGCGTGGGGAGTG	GGGAGCCCCCACCAT
	GACAAGCATGGCCTG	CAGCGTGGGGAGTGC	GGAGCCCCCACCATC
	ACAAGCATGGCCTGT	AGCGTGGGGAGTGCT	GAGCCCCCACCATCC
	CAAGCATGGCCTGTA	GCGTGGGGAGTGCTG	AGCCCCCACCATCCG
10	AAGCATGGCCTGTAC	CGTGGGGAGTGCTGG	GCCCCCACCATCCGG
	AGCATGGCCTGTACA	GTGGGGAGTGCTGGT	CCCCCACCATCCGGG
	GCATGGCCTGTACAA	TGGGGAGTGCTGGTG	CCCCACCATCCGGGG
	CATGGCCTGTACAAC	GGGGAGTGCTGGTGT	CCCACCATCCGGGGG
	ATGGCCTGTACAACC	GGGAGTGCTGGTGTG	CCACCATCCGGGGGG
15	TGGCCTGTACAACCT	GGAGTGCTGGTGTGT	CACCATCCGGGGGGA
	GGCCTGTACAACCTC	GAGTGCTGGTGTGTG	ACCATCCGGGGGGAC
	GCCTGTACAACCTCA	AGTGCTGGTGTGTGA	CCATCCGGGGGGACC
	CCTGTACAACCTCAA	GTGCTGGTGTGTGAA	CATCCGGGGGGACCC
	CTGTACAACCTCAAA	TGCTGGTGTGTGAAC	ATCCGGGGGGACCCC
20	TGTACAACCTCAAAC	GCTGGTGTGTGAACC	TCCGGGGGGACCCCG
	GTACAACCTCAAACA	CTGGTGTGTGAACCC	CCGGGGGGACCCCGA
	TACAACCTCAAACAG	TGGTGTGTGAACCCC	CGGGGGGACCCCGAG
	ACAACCTCAAACAGT	GGTGTGTGAACCCCA	GGGGGGACCCCGAGT
	CAACCTCAAACAGTG	GTGTGTGAACCCCAA	GGGGGACCCCGAGTG
25	AACCTCAAACAGTGC	TGTGTGAACCCCAAC	GGGGACCCCGAGTGT
	ACCTCAAACAGTGCA	GTGTGAACCCCAACA	GGGACCCCGAGTGTC
	CCTCAAACAGTGCAA	TGTGAACCCCAACAC	GGACCCCGAGTGTCA
	CTCAAACAGTGCAAG	GTGAACCCCAACACC	GACCCCGAGTGTCAT
	TCAAACAGTGCAAGA	TGAACCCCAACACCG	ACCCCGAGTGTCATC
30	CAAACAGTGCAAGAT	GAACCCCAACACCGG	CCCCGAGTGTCATCT
	AAACAGTGCAAGATG	AACCCCAACACCGGG	CCCGAGTGTCATCTC
	AACAGTGCAAGATGT	ACCCCAACACCGGGA	CCGAGTGTCATCTCT
	ACAGTGCAAGATGTC	CCCCAACACCGGGAA	CGAGTGTCATCTCTT
	CAGTGCAAGATGTCT	CCCAACACCGGGAAG	GAGTGTCATCTCTTC
35	AGTGCAAGATGTCTC	CCAACACCGGGAAGC	AGTGTCATCTCTTCT
	GTGCAAGATGTCTCT	CAACACCGGGAAGCT	GTGTCATCTCTTCTA
	TGCAAGATGTCTCTG	AACACCGGGAAGCTG	TGTCATCTCTTCTAC
	GCAAGATGTCTCTGA	ACACCGGGAAGCTGA	GTCATCTCTTCTACA
	CAAGATGTCTCTGAA	CACCGGGAAGCTGAT	TCATCTCTTCTACAA
40	AAGATGTCTCTGAAC	ACCGGGAAGCTGATC	CATCTCTTCTACAAT
	AGATGTCTCTGAACG	CCGGGAAGCTGATCC	ATCTCTTCTACAATG
	GATGTCTCTGAACGG	CGGGAAGCTGATCCA	TCTCTTCTACAATGA
	ATGTCTCTGAACGGG	GGGAAGCTGATCCAG	CTCTTCTACAATGAG
	TGTCTCTGAACGGGC	GGAAGCTGATCCAGG	TCTTCTACAATGAGC
45	GTCTCTGAACGGGCA	GAAGCTGATCCAGGG	CTTCTACAATGAGCA
	TCTCTGAACGGGCAG	AAGCTGATCCAGGGA	TTCTACAATGAGCAG
	CTCTGAACGGGCAGC	AGCTGATCCAGGGAG	TCTACAATGAGCAGC
	TCTGAACGGGCAGCG	GCTGATCCAGGGAGC	CTACAATGAGCAGCA
	CTGAACGGGCAGCGT	CTGATCCAGGGAGCC	TACAATGAGCAGCAG
50	TGAACGGGCAGCGTG	TGATCCAGGGAGCCC	ACAATGAGCAGCAGG
	GAACGGGCAGCGTGG	GATCCAGGGAGCCCC	CAATGAGCAGCAGGA

	AATGAGCAGCAGGAG	GCAGCCAGCCGGTGC	GCAGAAAACGGAGAG
	ATGAGCAGCAGGAGG	CAGCCAGCCGGTGCC	CAGAAAACGGAGAGT
	TGAGCAGCAGGAGGC	AGCCAGCCGGTGCCT	AGAAAACGGAGAGTG
	GAGCAGCAGGAGGCT	GCCAGCCGGTGCCTG	GAAAACGGAGAGTGC
5	AGCAGCAGGAGGCTT	CCAGCCGGTGCCTGG	AAAACGGAGAGTGCT
	GCAGCAGGAGGCTTG	CAGCCGGTGCCTGGC	AAACGGAGAGTGCTT
	CAGCAGGAGGCTTGC	AGCCGGTGCCTGGCG	AACGGAGAGTGCTTG
	AGCAGGAGGCTTGCG	GCCGGTGCCTGGCGC	ACGGAGAGTGCTTGG
	GCAGGAGGCTTGCGG	CCGGTGCCTGGCGCC	CGGAGAGTGCTTGGG
10	CAGGAGGCTTGCGGG	CGGTGCCTGGCGCCC	GGAGAGTGCTTGGGT
	AGGAGGCTTGCGGGG	GGTGCCTGGCGCCCC	GAGAGTGCTTGGGTG
	GGAGGCTTGCGGGGT	GTGCCTGGCGCCCCT	AGAGTGCTTGGGTGG
	GAGGCTTGCGGGGTG	TGCCTGGCGCCCCTG	GAGTGCTTGGGTGGT
	AGGCTTGCGGGGTGC	GCCTGGCGCCCCTGC	AGTGCTTGGGTGGTG
15	GGCTTGCGGGGTGCA	CCTGGCGCCCCTGCC	GTGCTTGGGTGGTGG
	GCTTGCGGGGTGCAC	CTGGCGCCCCTGCCC	TGCTTGGGTGGTGGG
	CTTGCGGGGTGCACA	TGGCGCCCCTGCCCC	GCTTGGGTGGTGGGT
	TTGCGGGGTGCACAC	GGCGCCCCTGCCCCC	CTTGGGTGGTGGGTG
	TGCGGGGTGCACACC	GCGCCCCTGCCCCCC	TTGGGTGGTGGGTGC
20	GCGGGGTGCACACCC	CGCCCCTGCCCCCCG	TGGGTGGTGGGTGCT
	CGGGGTGCACACCCA	GCCCCTGCCCCCCGC	GGGTGGTGGGTGCTG
	GGGGTGCACACCCAG	CCCCTGCCCCCCGCC	GGTGGTGGGTGCTGG
	GGGTGCACACCCAGC	CCCTGCCCCCCGCCC	GTGGTGGGTGCTGGA
	GGTGCACACCCAGCG	CCTGCCCCCCGCCCC	TGGTGGGTGCTGGAG
25	GTGCACACCCAGCGG	CTGCCCCCCGCCCCT	GGTGGGTGCTGGAGG
	TGCACACCCAGCGGA	TGCCCCCCGCCCCTC	GTGGGTGCTGGAGGA
	GCACACCCAGCGGAT	GCCCCCCGCCCCTCT	TGGGTGCTGGAGGAT
	CACACCCAGCGGATG	CCCCCCGCCCCTCTC	GGGTGCTGGAGGATT
	ACACCCAGCGGATGC	CCCCCGCCCCTCTCC	GGTGCTGGAGGATTT
30	CACCCAGCGGATGCA	CCCCGCCCCTCTCCA	GTGCTGGAGGATTTT
	ACCCAGCGGATGCAG	CCCGCCCCTCTCCAA	TGCTGGAGGATTTTC
	CCCAGCGGATGCAGT	CCGCCCCTCTCCAAA	GCTGGAGGATTTTCC
	CCAGCGGATGCAGTA	CGCCCCTCTCCAAAC	CTGGAGGATTTTCCA
	CAGCGGATGCAGTAG	GCCCCTCTCCAAACA	TGGAGGATTTTCCAG
35	AGCGGATGCAGTAGA	CCCCTCTCCAAACAC	GGAGGATTTTCCAGT
	GCGGATGCAGTAGAC	CCCTCTCCAAACACC	GAGGATTTTCCAGTT
	CGGATGCAGTAGACC	CCTCTCCAAACACCG	AGGATTTTCCAGTTC
	GGATGCAGTAGACCG	CTCTCCAAACACCGG	GGATTTTCCAGTTCT
	GATGCAGTAGACCGC	TCTCCAAACACCGGC	GATTTTCCAGTTCTG
40	ATGCAGTAGACCGCA	CTCCAAACACCGGCA	ATTTTCCAGTTCTGA
	TGCAGTAGACCGCAG	TCCAAACACCGGCAG	TTTTCCAGTTCTGAC
	GCAGTAGACCGCAGC	CCAAACACCGGCAGA	TTTCCAGTTCTGACA
	CAGTAGACCGCAGCC	CAAACACCGGCAGAA	TTCCAGTTCTGACAC
	AGTAGACCGCAGCCA	AAACACCGGCAGAAA	TCCAGTTCTGACACA
45	GTAGACCGCAGCCAG	AACACCGGCAGAAAA	CCAGTTCTGACACAC
	TAGACCGCAGCCAGC	ACACCGGCAGAAAAC	CAGTTCTGACACACG
	AGACCGCAGCCAGCC	CACCGGCAGAAAACG	AGTTCTGACACACGT
	GACCGCAGCCAGCCG	ACCGGCAGAAAACGG	GTTCTGACACACGTA
	ACCGCAGCCAGCCGG	CCGGCAGAAAACGGA	TTCTGACACACGTAT
50	CCGCAGCCAGCCGGT	CGGCAGAAAACGGAG	TCTGACACACGTATT
	CGCAGCCAGCCGGTG	GGCAGAAAACGGAGA	CTGACACACGTATTT

TGACACACGTATTTA	CCCGGCCTCTCTCTT	TCCCCGGGGGAGGAA
GACACACGTATTTAT	CCGGCCTCTCTCTTC	CCCCGGGGGAGGAAG
ACACACGTATTTATA	CGGCCTCTCTCTTCC	CCCGGGGGAGGAAGG
CACACGTATTTATAT	GGCCTCTCTCTTCCC	CCGGGGGAGGAAGGG
ACACGTATTTATATT	GCCTCTCTCTTCCCA	CGGGGGAGGAAGGGG
CACGTATTTATATTT	CCTCTCTCTTCCCAG	GGGGGAGGAAGGGGG
ACGTATTTATATTTG	CTCTCTCTTCCCAGC	GGGGAGGAAGGGGGT
CGTATTTATATTTGG	TCTCTCTTCCCAGCT	GGGAGGAAGGGGGTT
GTATTTATATTTGGA	CTCTCTTCCCAGCTG	GGAGGAAGGGGGTTG
TATTTATATTTGGAA	TCTCTTCCCAGCTGC	GAGGAAGGGGGTTGT
ATTTATATTTGGAAA	CTCTTCCCAGCTGCA	AGGAAGGGGGTTGTG
TTTATATTTGGAAAG	TCTTCCCAGCTGCAG	GGAAGGGGGTTGTGG
TTATATTTGGAAAGA	CTTCCCAGCTGCAGA	GAAGGGGGTTGTGGT
TATATTTGGAAAGAG	TTCCCAGCTGCAGAT	AAGGGGGTTGTGGTC
ATATTTGGAAAGAGA	TCCCAGCTGCAGATG	AGGGGGTTGTGGTCG
TATTTGGAAAGAGAC	CCCAGCTGCAGATGC	GGGGGTTGTGGTCGG
ATTTGGAAAGAGACC	CCAGCTGCAGATGCC	GGGGTTGTGGTCGGG
TTTGGAAAGAGACCA	CAGCTGCAGATGCCA	GGGTTGTGGTCGGGG
TTGGAAAGAGACCAG	AGCTGCAGATGCCAC	GGTTGTGGTCGGGGA
TGGAAAGAGACCAGC	GCTGCAGATGCCACA	GTTGTGGTCGGGGAG
GGAAAGAGACCAGCA	CTGCAGATGCCACAC	TTGTGGTCGGGGAGC
GAAAGAGACCAGCAC	TGCAGATGCCACACC	TGTGGTCGGGGAGCT
AAAGAGACCAGCACC	GCAGATGCCACACCT	GTGGTCGGGGAGCTG
AAGAGACCAGCACCG	CAGATGCCACACCTG	TGGTCGGGGAGCTGG
AGAGACCAGCACCGA	AGATGCCACACCTGC	GGTCGGGGAGCTGGG
GAGACCAGCACCGAG	GATGCCACACCTGCT	GTCGGGGAGCTGGGG
AGACCAGCACCGAGC	ATGCCACACCTGCTC	TCGGGGAGCTGGGGT
GACCAGCACCGAGCT	TGCCACACCTGCTCC	CGGGGAGCTGGGGTA
ACCAGCACCGAGCTC	GCCACACCTGCTCCT	GGGGAGCTGGGGTAC
CCAGCACCGAGCTCG	CCACACCTGCTCCTT	GGGAGCTGGGGTACA
CAGCACCGAGCTCGG	CACACCTGCTCCTTC	GGAGCTGGGGTACAG
AGCACCGAGCTCGGC	ACACCTGCTCCTTCT	GAGCTGGGGTACAGG
GCACCGAGCTCGGCA	CACCTGCTCCTTCTT	AGCTGGGGTACAGGT
CACCGAGCTCGGCAC	ACCTGCTCCTTCTTG	GCTGGGGTACAGGTT
ACCGAGCTCGGCACC	CCTGCTCCTTCTTGC	CTGGGGTACAGGTTT
CCGAGCTCGGCACCT	CTGCTCCTTCTTGCT	TGGGGTACAGGTTTG
CGAGCTCGGCACCTC	TGCTCCTTCTTGCTT	GGGGTACAGGTTTGG
GAGCTCGGCACCTCC	GCTCCTTCTTGCTTT	GGGTACAGGTTTGGG
AGCTCGGCACCTCCC	CTCCTTCTTGCTTTC	GGTACAGGTTTGGGG
GCTCGGCACCTCCCC	TCCTTCTTGCTTTCC	GTACAGGTTTGGGGA
CTCGGCACCTCCCCG	CCTTCTTGCTTTCCC	TACAGGTTTGGGGAG
TCGGCACCTCCCCGG	CTTCTTGCTTTCCCC	ACAGGTTTGGGGAGG
CGGCACCTCCCCGGC	TTCTTGCTTTCCCCG	CAGGTTTGGGGAGGG
GGCACCTCCCCGGCC	TCTTGCTTTCCCCGG	AGGTTTGGGGAGGGG
GCACCTCCCCGGCCT	CTTGCTTTCCCCGGG	GGTTTGGGGAGGGGG
CACCTCCCCGGCCTC	TTGCTTTCCCCGGGG	GTTTGGGGAGGGGGA
ACCTCCCCGGCCTCT	TGCTTTCCCCGGGGG	TTTGGGGAGGGGGAA
CCTCCCCGGCCTCTC	GCTTTCCCCGGGGGA	TTGGGGAGGGGGAAG
CTCCCCGGCCTCTCT	CTTTCCCCGGGGGAG	TGGGGAGGGGGAAGA
TCCCCGGCCTCTCTC	TTTCCCCGGGGGAGG	GGGGAGGGGGAAGAG
CCCCGGCCTCTCTCT	TTCCCCGGGGGAGGA	GGGAGGGGGAAGAGA

GGAGGGGGAAGAGAA	AGATTAAAGGAAGGA
GAGGGGGAAGAGAAA	GATTAAAGGAAGGAA
AGGGGGAAGAGAAAT	ATTAAAGGAAGGAAA
GGGGGAAGAGAAATT	TTAAAGGAAGGAAAA
GGGGAAGAGAAATTT	TAAAGGAAGGAAAAG
GGGAAGAGAAATTTT	AAAGGAAGGAAAAGT
GGAAGAGAAATTTTT	
GAAGAGAAATTTTTA	
AAGAGAAATTTTTAT	
AGAGAAATTTTTATT	
GAGAAATTTTTATTT	
AGAAATTTTTATTTT	
GAAATTTTTATTTTT	
AAATTTTTATTTTTG	
AATTTTTATTTTTGA	
ATTTTTATTTTTGAA	
TTTTTATTTTTGAAC	
TTTTATTTTTGAACC	
TTTATTTTTGAACCC	
TTATTTTTGAACCCC	
TATTTTTGAACCCCT	
ATTTTTGAACCCCTG	
TTTTTGAACCCCTGT	
TTTTGAACCCCTGTG	
TTTGAACCCCTGTGT	
TTGAACCCCTGTGTC	
TGAACCCCTGTGTCC	
GAACCCCTGTGTCCC	
AACCCCTGTGTCCCT	
ACCCCTGTGTCCCTT	
CCCCTGTGTCCCTTT	
CCCTGTGTCCCTTTT	
CCTGTGTCCCTTTTG	
CTGTGTCCCTTTTGC	
TGTGTCCCTTTTGCA	
GTGTCCCTTTTGCAT	
TGTCCCTTTTGCATA	
GTCCCTTTTGCATAA	
TCCCTTTTGCATAAG	
CCCTTTTGCATAAGA	
CCTTTTGCATAAGAT	
CTTTTGCATAAGATT	
TTTTGCATAAGATTA	
TTTGCATAAGATTAA	
TTGCATAAGATTAAA	
TGCATAAGATTAAAG	
GCATAAGATTAAAGG	
CATAAGATTAAAGGA	
ATAAGATTAAAGGAA	
TAAGATTAAAGGAAG	
AAGATTAAAGGAAGG	

EXAMPLE 7

Antisense oligonucleotides to IGFBP3 may be selected from molecules capable of interacting with one or more of the following sense oligonucleotides:

CTCAGCGCCCAGCCG	GCCGTGTACTGTCGC	GCAGCGTGCCCCGGT
TCAGCGCCCAGCCGC	CCGTGTACTGTCGCC	CAGCGTGCCCCGGTT
CAGCGCCCAGCCGCT	CGTGTACTGTCGCCC	AGCGTGCCCCGGTTG
AGCGCCCAGCCGCTT	GTGTACTGTCGCCCC	GCGTGCCCCGGTTGC
GCGCCCAGCCGCTTC	TGTACTGTCGCCCCA	CGTGCCCCGGTTGCA
CGCCCAGCCGCTTCC	GTACTGTCGCCCCAT	GTGCCCCGGTTGCAG
GCCCAGCCGCTTCCT	TACTGTCGCCCCATC	TGCCCCGGTTGCAGG
CCCAGCCGCTTCCTG	ACTGTCGCCCCATCC	GCCCCGGTTGCAGGC
CCAGCCGCTTCCTGC	CTGTCGCCCCATCCC	CCCCGGTTGCAGGCG
CAGCCGCTTCCTGCC	TGTCGCCCCATCCCT	CCCGGTTGCAGGCGT
AGCCGCTTCCTGCCT	GTCGCCCCATCCCTG	CCGGTTGCAGGCGTC
GCCGCTTCCTGCCTG	TCGCCCCATCCCTGC	CGGTTGCAGGCGTCA
CCGCTTCCTGCCTGG	CGCCCCATCCCTGCG	GGTTGCAGGCGTCAT
CGCTTCCTGCCTGGA	GCCCCATCCCTGCGC	GTTGCAGGCGTCATG
GCTTCCTGCCTGGAT	CCCCATCCCTGCGCG	TTGCAGGCGTCATGC
CTTCCTGCCTGGATT	CCCATCCCTGCGCGC	TGCAGGCGTCATGCA
TTCCTGCCTGGATTC	CCATCCCTGCGCGCC	GCAGGCGTCATGCAG
TCCTGCCTGGATTCC	CATCCCTGCGCGCCC	CAGGCGTCATGCAGC
CCTGCCTGGATTCCA	ATCCCTGCGCGCCCA	AGGCGTCATGCAGCG
CTGCCTGGATTCCAC	TCCCTGCGCGCCCAG	GGCGTCATGCAGCGG
TGCCTGGATTCCACA	CCCTGCGCGCCCAGC	GCGTCATGCAGCGGG
GCCTGGATTCCACAG	CCTGCGCGCCCAGCC	CGTCATGCAGCGGGC
CCTGGATTCCACAGC	CTGCGCGCCCAGCCT	GTCATGCAGCGGGCG
CTGGATTCCACAGCT	TGCGCGCCCAGCCTG	TCATGCAGCGGGCGC
TGGATTCCACAGCTT	GCGCGCCCAGCCTGC	CATGCAGCGGGCGCG
GGATTCCACAGCTTC	CGCGCCCAGCCTGCC	ATGCAGCGGGCGCGA
GATTCCACAGCTTCG	GCGCCCAGCCTGCCA	TGCAGCGGGCGCGAC
ATTCCACAGCTTCGC	CGCCCAGCCTGCCAA	GCAGCGGGCGCGACC
TTCCACAGCTTCGCG	GCCCAGCCTGCCAAG	CAGCGGGCGCGACCC
TCCACAGCTTCGCGC	CCCAGCCTGCCAAGC	AGCGGGCGCGACCCA
CCACAGCTTCGCGCC	CCAGCCTGCCAAGCA	GCGGGCGCGACCCAC
CACAGCTTCGCGCCG	CAGCCTGCCAAGCAG	CGGGCGCGACCCACG
ACAGCTTCGCGCCGT	AGCCTGCCAAGCAGC	GGGCGCGACCCACGC
CAGCTTCGCGCCGTG	GCCTGCCAAGCAGCG	GGCGCGACCCACGCT
AGCTTCGCGCCGTGT	CCTGCCAAGCAGCGT	GCGCGACCCACGCTC
GCTTCGCGCCGTGTA	CTGCCAAGCAGCGTG	CGCGACCCACGCTCT
CTTCGCGCCGTGTAC	TGCCAAGCAGCGTGC	GCGACCCACGCTCTG
TTCGCGCCGTGTACT	GCCAAGCAGCGTGCC	CGACCCACGCTCTGG
TCGCGCCGTGTACTG	CCAAGCAGCGTGCCC	GACCCACGCTCTGGG
CGCGCCGTGTACTGT	CAAGCAGCGTGCCCC	ACCCACGCTCTGGGC
GCGCCGTGTACTGTC	AAGCAGCGTGCCCCG	CCCACGCTCTGGGCC
CGCCGTGTACTGTCG	AGCAGCGTGCCCCGG	CCACGCTCTGGGCCG

CACGCTCTGGGCCGC	GGTGGCGCGGGCTGG	CGAGCCGTGCGACGC
ACGCTCTGGGCCGCT	GTGGCGCGGGCTGGC	GAGCCGTGCGACGCG
CGCTCTGGGCCGCTG	TGGCGCGGGCTGGCG	AGCCGTGCGACGCGC
GCTCTGGGCCGCTGC	GGCGCGGGCTGGCGC	GCCGTGCGACGCGCG
CTCTGGGCCGCTGCG	GCGCGGGCTGGCGCG	CCGTGCGACGCGCGT
TCTGGGCCGCTGCGC	CGCGGGCTGGCGCGA	CGTGCGACGCGCGTG
CTGGGCCGCTGCGCT	GCGGGCTGGCGCGAG	GTGCGACGCGCGTGC
TGGGCCGCTGCGCTG	CGGGCTGGCGCGAGC	TGCGACGCGCGTGCA
GGGCCGCTGCGCTGA	GGGCTGGCGCGAGCT	GCGACGCGCGTGCAC
GGCCGCTGCGCTGAC	GGCTGGCGCGAGCTC	CGACGCGCGTGCACT
GCCGCTGCGCTGACT	GCTGGCGCGAGCTCG	GACGCGCGTGCACTG
CCGCTGCGCTGACTC	CTGGCGCGAGCTCGG	ACGCGCGTGCACTGG
CGCTGCGCTGACTCT	TGGCGCGAGCTCGGG	CGCGCGTGCACTGGC
GCTGCGCTGACTCTG	GGCGCGAGCTCGGGG	GCGCGTGCACTGGCC
CTGCGCTGACTCTGC	GCGCGAGCTCGGGGG	CGCGTGCACTGGCCC
TGCGCTGACTCTGCT	CGCGAGCTCGGGGGG	GCGTGCACTGGCCCA
GCGCTGACTCTGCTG	GCGAGCTCGGGGGGC	CGTGCACTGGCCCAG
CGCTGACTCTGCTGG	CGAGCTCGGGGGGCT	GTGCACTGGCCCAGT
GCTGACTCTGCTGGT	GAGCTCGGGGGGCTT	TGCACTGGCCCAGTG
CTGACTCTGCTGGTG	AGCTCGGGGGGCTTG	GCACTGGCCCAGTGC
TGACTCTGCTGGTGC	GCTCGGGGGGCTTGG	CACTGGCCCAGTGCG
GACTCTGCTGGTGCT	CTCGGGGGGCTTGGG	ACTGGCCCAGTGCGC
ACTCTGCTGGTGCTG	TCGGGGGGCTTGGGT	CTGGCCCAGTGCGCG
CTCTGCTGGTGCTGC	CGGGGGGCTTGGGTC	TGGCCCAGTGCGCGC
TCTGCTGGTGCTGCT	GGGGGGCTTGGGTCC	GGCCCAGTGCGCGCC
CTGCTGGTGCTGCTC	GGGGGCTTGGGTCCC	GCCCAGTGCGCGCCT
TGCTGGTGCTGCTCC	GGGGCTTGGGTCCCG	CCCAGTGCGCGCCTC
GCTGGTGCTGCTCCG	GGGCTTGGGTCCCGT	CCAGTGCGCGCCTCC
CTGGTGCTGCTCCGC	GGCTTGGGTCCCGTG	CAGTGCGCGCCTCCG
TGGTGCTGCTCCGCG	GCTTGGGTCCCGTGG	AGTGCGCGCCTCCGC
GGTGCTGCTCCGCGG	CTTGGGTCCCGTGGT	GTGCGCGCCTCCGCC
GTGCTGCTCCGCGGG	TTGGGTCCCGTGGTG	TGCGCGCCTCCGCCC
TGCTGCTCCGCGGGC	TGGGTCCCGTGGTGC	GCGCGCCTCCGCCCG
GCTGCTCCGCGGGCC	GGGTCCCGTGGTGCG	CGCGCCTCCGCCCGC
CTGCTCCGCGGGCCG	GGTCCCGTGGTGCGC	GCGCCTCCGCCCGCC
TGCTCCGCGGGCCGC	GTCCCGTGGTGCGCT	CGCCTCCGCCCGCCG
GCTCCGCGGGCCGCC	TCCCGTGGTGCGCTG	GCCTCCGCCCGCCGT
CTCCGCGGGCCGCCG	CCCGTGGTGCGCTGC	CCTCCGCCCGCCGTG
TCCGCGGGCCGCCGG	CCGTGGTGCGCTGCG	CTCCGCCCGCCGTGT
CCGCGGGCCGCCGGT	CGTGGTGCGCTGCGA	TCCGCCCGCCGTGTG
CGCGGGCCGCCGGTG	GTGGTGCGCTGCGAG	CCGCCCGCCGTGTGC
GCGGGCCGCCGGTGG	TGGTGCGCTGCGAGC	CGCCCGCCGTGTGCG
CGGGCCGCCGGTGGC	GGTGCGCTGCGAGCC	GCCCGCCGTGTGCGC
GGGCCGCCGGTGGCG	GTGCGCTGCGAGCCG	CCCGCCGTGTGCGCG
GGCCGCCGGTGGCGC	TGCGCTGCGAGCCGT	CCGCCGTGTGCGCGG
GCCGCCGGTGGCGCG	GCGCTGCGAGCCGTG	CGCCGTGTGCGCGGA
CCGCCGGTGGCGCGG	CGCTGCGAGCCGTGC	GCCGTGTGCGCGGAG
CGCCGGTGGCGCGGG	GCTGCGAGCCGTGCG	CCGTGTGCGCGGAGC
GCCGGTGGCGCGGGC	CTGCGAGCCGTGCGA	CGTGTGCGCGGAGCT
CCGGTGGCGCGGGCT	TGCGAGCCGTGCGAC	GTGTGCGCGGAGCTG
CGGTGGCGCGGGCTG	GCGAGCCGTGCGACG	TGTGCGCGGAGCTGG

	GTGCGCGGAGCTGGT	ACTGAGCGAGGGCCA	CCTTCGCTGCCAGCC
	TGCGCGGAGCTGGTG	CTGAGCGAGGGCCAG	CTTCGCTGCCAGCCG
	GCGCGGAGCTGGTGC	TGAGCGAGGGCCAGC	TTCGCTGCCAGCCGT
	CGCGGAGCTGGTGCG	GAGCGAGGGCCAGCC	TCGCTGCCAGCCGTC
5	GCGGAGCTGGTGCGC	AGCGAGGGCCAGCCG	CGCTGCCAGCCGTCG
	CGGAGCTGGTGCGCG	GCGAGGGCCAGCCGT	GCTGCCAGCCGTCGC
	GGAGCTGGTGCGCGA	CGAGGGCCAGCCGTG	CTGCCAGCCGTCGCC
	GAGCTGGTGCGCGAG	GAGGGCCAGCCGTGC	TGCCAGCCGTCGCCC
	AGCTGGTGCGCGAGC	AGGGCCAGCCGTGCG	GCCAGCCGTCGCCCG
10	GCTGGTGCGCGAGCC	GGGCCAGCCGTGCGG	CCAGCCGTCGCCCGA
	CTGGTGCGCGAGCCG	GGCCAGCCGTGCGGC	CAGCCGTCGCCCGAC
	TGGTGCGCGAGCCGG	GCCAGCCGTGCGGCA	AGCCGTCGCCCGACG
	GGTGCGCGAGCCGGG	CCAGCCGTGCGGCAT	GCCGTCGCCCGACGA
	GTGCGCGAGCCGGGC	CAGCCGTGCGGCATC	CCGTCGCCCGACGAG
15	TGCGCGAGCCGGGCT	AGCCGTGCGGCATCT	CGTCGCCCGACGAGG
	GCGCGAGCCGGGCTG	GCCGTGCGGCATCTA	GTCGCCCGACGAGGC
	CGCGAGCCGGGCTGC	CCGTGCGGCATCTAC	TCGCCCGACGAGGCG
	GCGAGCCGGGCTGCG	CGTGCGGCATCTACA	CGCCCGACGAGGCGC
	CGAGCCGGGCTGCGG	GTGCGGCATCTACAC	GCCCGACGAGGCGCG
20	GAGCCGGGCTGCGGC	TGCGGCATCTACACC	CCCGACGAGGCGCGA
	AGCCGGGCTGCGGCT	GCGGCATCTACACCG	CCGACGAGGCGCGAC
	GCCGGGCTGCGGCTG	CGGCATCTACACCGA	CGACGAGGCGCGACC
	CCGGGCTGCGGCTGC	GGCATCTACACCGAG	GACGAGGCGCGACCG
	CGGGCTGCGGCTGCT	GCATCTACACCGAGC	ACGAGGCGCGACCGC
25	GGGCTGCGGCTGCTG	CATCTACACCGAGCG	CGAGGCGCGACCGCT
	GGCTGCGGCTGCTGC	ATCTACACCGAGCGC	GAGGCGCGACCGCTG
	GCTGCGGCTGCTGCC	TCTACACCGAGCGCT	AGGCGCGACCGCTGC
	CTGCGGCTGCTGCCT	CTACACCGAGCGCTG	GGCGCGACCGCTGCA
	TGCGGCTGCTGCCTG	TACACCGAGCGCTGT	GCGCGACCGCTGCAG
30	GCGGCTGCTGCCTGA	ACACCGAGCGCTGTG	CGCGACCGCTGCAGG
	CGGCTGCTGCCTGAC	CACCGAGCGCTGTGG	GCGACCGCTGCAGGC
	GGCTGCTGCCTGACG	ACCGAGCGCTGTGGC	CGACCGCTGCAGGCG
	GCTGCTGCCTGACGT	CCGAGCGCTGTGGCT	GACCGCTGCAGGCGC
	CTGCTGCCTGACGTG	CGAGCGCTGTGGCTC	ACCGCTGCAGGCGCT
35	TGCTGCCTGACGTGC	GAGCGCTGTGGCTCC	CCGCTGCAGGCGCTG
	GCTGCCTGACGTGCG	AGCGCTGTGGCTCCG	CGCTGCAGGCGCTGC
	CTGCCTGACGTGCGC	GCGCTGTGGCTCCGG	GCTGCAGGCGCTGCT
	TGCCTGACGTGCGCA	CGCTGTGGCTCCGGC	CTGCAGGCGCTGCTG
	GCCTGACGTGCGCAC	GCTGTGGCTCCGGCC	TGCAGGCGCTGCTGG
40	CCTGACGTGCGCACT	CTGTGGCTCCGGCCT	GCAGGCGCTGCTGGA
	CTGACGTGCGCACTG	TGTGGCTCCGGCCTT	CAGGCGCTGCTGGAC
	TGACGTGCGCACTGA	GTGGCTCCGGCCTTC	AGGCGCTGCTGGACG
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GAGACTCTGCCCTGC	CTCCTTGAAAACAGA	CTTTATTTTTTTAAC
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GACTCTGCCCTGCTG	CCTTGAAAACAGAGG	TTATTTTTTTAACTT
ACTCTGCCCTGCTGC	CTTGAAAACAGAGGG	TATTTTTTTAACTTT
CTCTGCCCTGCTGCA	TTGAAAACAGAGGGG	ATTTTTTTAACTTTT
TCTGCCCTGCTGCAG	TGAAAACAGAGGGGT	TTTTTTTAACTTTTT
CTGCCCTGCTGCAGA	GAAAACAGAGGGGTC	TTTTTTAACTTTTTG
TGCCCTGCTGCAGAC	AAAACAGAGGGGTCT	TTTTTAACTTTTTGG
GCCCTGCTGCAGACC	AAACAGAGGGGTCTC	TTTTAACTTTTTGGG
CCCTGCTGCAGACCT	AACAGAGGGGTCTCA	TTTAACTTTTTGGGG
CCTGCTGCAGACCTC	ACAGAGGGGTCTCAA	TTAACTTTTTGGGGG
CTGCTGCAGACCTCG	CAGAGGGGTCTCAAG	TAACTTTTTGGGGGG
TGCTGCAGACCTCGG	AGAGGGGTCTCAAGA	AACTTTTTGGGGGGA
GCTGCAGACCTCGGT	GAGGGGTCTCAAGAC	ACTTTTTGGGGGGAA
CTGCAGACCTCGGTG	AGGGGTCTCAAGACA	CTTTTTGGGGGGAAA
TGCAGACCTCGGTGT	GGGGTCTCAAGACAT	TTTTTGGGGGGAAAA
GCAGACCTCGGTGTG	GGGTCTCAAGACATT	TTTTGGGGGGAAAAG
CAGACCTCGGTGTGG	GGTCTCAAGACATTC	TTTGGGGGGAAAAGT
AGACCTCGGTGTGGA	GTCTCAAGACATTCT	TTGGGGGGAAAAGTA
GACCTCGGTGTGGAC	TCTCAAGACATTCTG	TGGGGGGAAAAGTAT
ACCTCGGTGTGGACA	CTCAAGACATTCTGC	GGGGGGAAAAGTATT
CCTCGGTGTGGACAC	TCAAGACATTCTGCC	GGGGGAAAAGTATTT
CTCGGTGTGGACACA	CAAGACATTCTGCCT	GGGGAAAAGTATTTT
TCGGTGTGGACACAC	AAGACATTCTGCCTA	GGGAAAAGTATTTTT
CGGTGTGGACACACG	AGACATTCTGCCTAC	GGAAAAGTATTTTTG
GGTGTGGACACACGC	GACATTCTGCCTACC	GAAAAGTATTTTTGA
GTGTGGACACACGCT	ACATTCTGCCTACCT	AAAAGTATTTTTGAG
TGTGGACACACGCTG	CATTCTGCCTACCTA	AAAGTATTTTTGAGA
GTGGACACACGCTGC	ATTCTGCCTACCTAT	AAGTATTTTTGAGAA
TGGACACACGCTGCA	TTCTGCCTACCTATT	AGTATTTTTGAGAAG
GGACACACGCTGCAT	TCTGCCTACCTATTA	GTATTTTTGAGAAGT
GACACACGCTGCATA	CTGCCTACCTATTAG	TATTTTTGAGAAGTT
ACACACGCTGCATAG	TGCCTACCTATTAGC	ATTTTTGAGAAGTTT
CACACGCTGCATAGA	GCCTACCTATTAGCT	TTTTTGAGAAGTTTG
ACACGCTGCATAGAG	CCTACCTATTAGCTT	TTTTGAGAAGTTTGT
CACGCTGCATAGAGC	CTACCTATTAGCTTT	TTTGAGAAGTTTGTC
ACGCTGCATAGAGCT	TACCTATTAGCTTTT	TTGAGAAGTTTGTCT
CGCTGCATAGAGCTC	ACCTATTAGCTTTTC	TGAGAAGTTTGTCTT

GAGAAGTTTGTCTTG
AGAAGTTTGTCTTGC
GAAGTTTGTCTTGCA
AAGTTTGTCTTGCAA
AGTTTGTCTTGCAAT
GTTTGTCTTGCAATG
TTTGTCTTGCAATGT
TTGTCTTGCAATGTA
TGTCTTGCAATGTAT
GTCTTGCAATGTATT
TCTTGCAATGTATTT
CTTGCAATGTATTTA
TTGCAATGTATTTAT
TGCAATGTATTTATA
GCAATGTATTTATAA
CAATGTATTTATAAA
AATGTATTTATAAAT
ATGTATTTATAAATA
TGTATTTATAAATAG
GTATTTATAAATAGT
TATTTATAAATAGTA
ATTTATAAATAGTAA
TTTATAAATAGTAAA
TTATAAATAGTAAAT
TATAAATAGTAAATA
ATAAATAGTAAATAA
TAAATAGTAAATAAA
AAATAGTAAATAAAG
AATAGTAAATAAAGT
ATAGTAAATAAAGTT
TAGTAAATAAAGTTT
AGTAAATAAAGTTTT
GTAAATAAAGTTTTT
TAAATAAAGTTTTTA
AAATAAAGTTTTTAC
AATAAAGTTTTTACC
ATAAAGTTTTTACCA
TAAAGTTTTTACCAT
AAAGTTTTTACCATT

EXAMPLE 8

Antisense oligonucleotides to IGF-I may be selected from molecules capable of interacting with one or more of the following sense oligonucleotides:

TTTTTTTTTTTTTTG	TTTTTTTTTTTGAGA	TTTTTTTTGAGAAAG
TTTTTTTTTTTTTGA	TTTTTTTTTTGAGAA	TTTTTTTGAGAAAGG
TTTTTTTTTTTTGAG	TTTTTTTTTGAGAAA	TTTTTTGAGAAAGGG

	TTTTTGAGAAAGGGA	GGAGGAGGGTCCCCG	CTCTCGCTCTGGCCG
	TTTTGAGAAAGGGAA	GAGGAGGGTCCCCGA	TCTCGCTCTGGCCGA
	TTTGAGAAAGGGAAT	AGGAGGGTCCCCGAC	CTCGCTCTGGCCGAC
	TTGAGAAAGGGAATT	GGAGGGTCCCCGACC	TCGCTCTGGCCGACG
5	TGAGAAAGGGAATTT	GAGGGTCCCCGACCT	CGCTCTGGCCGACGA
	GAGAAAGGGAATTTC	AGGGTCCCCGACCTC	GCTCTGGCCGACGAG
	AGAAAGGGAATTTCA	GGGTCCCCGACCTCG	CTCTGGCCGACGAGT
	GAAAGGGAATTTCAT	GGTCCCCGACCTCGC	TCTGGCCGACGAGTG
	AAAGGGAATTTCATC	GTCCCCGACCTCGCT	CTGGCCGACGAGTGG
10	AAGGGAATTTCATCC	TCCCCGACCTCGCTG	TGGCCGACGAGTGGA
	AGGGAATTTCATCCC	CCCCGACCTCGCTGT	GGCCGACGAGTGGAG
	GGGAATTTCATCCCA	CCCGACCTCGCTGTG	GCCGACGAGTGGAGA
	GGAATTTCATCCCAA	CCGACCTCGCTGTGG	CCGACGAGTGGAGAA
	GAATTTCATCCCAAA	CGACCTCGCTGTGGG	CGACGAGTGGAGAAA
15	AATTTCATCCCAAAT	GACCTCGCTGTGGGG	GACGAGTGGAGAAAT
	ATTTCATCCCAAATA	ACCTCGCTGTGGGGG	ACGAGTGGAGAAATC
	TTTCATCCCAAATAA	CCTCGCTGTGGGGGC	CGAGTGGAGAAATCT
	TTCATCCCAAATAAA	CTCGCTGTGGGGGCT	GAGTGGAGAAATCTG
	TCATCCCAAATAAAA	TCGCTGTGGGGGCTC	AGTGGAGAAATCTGC
20	CATCCCAAATAAAAG	CGCTGTGGGGGCTCC	GTGGAGAAATCTGCG
	ATCCCAAATAAAAGG	GCTGTGGGGGCTCCT	TGGAGAAATCTGCGG
	TCCCAAATAAAAGGA	CTGTGGGGGCTCCTG	GGAGAAATCTGCGGG
	CCCAAATAAAAGGAA	TGTGGGGGCTCCTGT	GAGAAATCTGCGGGC
	CCAAATAAAAGGAAT	GTGGGGGCTCCTGTT	AGAAATCTGCGGGCC
25	CAAATAAAAGGAATG	TGGGGGCTCCTGTTT	GAAATCTGCGGGCCA
	AAATAAAAGGAATGA	GGGGGCTCCTGTTTC	AAATCTGCGGGCCAG
	AATAAAAGGAATGAA	GGGGCTCCTGTTTCT	AATCTGCGGGCCAGG
	ATAAAAGGAATGAAG	GGGCTCCTGTTTCTC	ATCTGCGGGCCAGGC
	TAAAAGGAATGAAGT	GGCTCCTGTTTCTCT	TCTGCGGGCCAGGCA
30	AAAAGGAATGAAGTC	GCTCCTGTTTCTCTC	CTGCGGGCCAGGCAT
	AAAGGAATGAAGTCT	CTCCTGTTTCTCTCC	TGCGGGCCAGGCATC
	AAGGAATGAAGTCTG	TCCTGTTTCTCTCCG	GCGGGCCAGGCATCG
	AGGAATGAAGTCTGG	CCTGTTTCTCTCCGC	CGGGCCAGGCATCGA
	GGAATGAAGTCTGGC	CTGTTTCTCTCCGCC	GGGCCAGGCATCGAC
35	GAATGAAGTCTGGCT	TGTTTCTCTCCGCCG	GGCCAGGCATCGACA
	AATGAAGTCTGGCTC	GTTTCTCTCCGCCGC	GCCAGGCATCGACAT
	ATGAAGTCTGGCTCC	TTTCTCTCCGCCGCG	CCAGGCATCGACATC
	TGAAGTCTGGCTCCG	TTCTCTCCGCCGCGC	CAGGCATCGACATCC
	GAAGTCTGGCTCCGG	TCTCTCCGCCGCGCT	AGGCATCGACATCCG
40	AAGTCTGGCTCCGGA	CTCTCCGCCGCGCTC	GGCATCGACATCCGC
	AGTCTGGCTCCGGAG	TCTCCGCCGCGCTCT	GCATCGACATCCGCA
	GTCTGGCTCCGGAGG	CTCCGCCGCGCTCTC	CATCGACATCCGCAA
	TCTGGCTCCGGAGGA	TCCGCCGCGCTCTCG	ATCGACATCCGCAAC
	CTGGCTCCGGAGGAG	CCGCCGCGCTCTCGC	TCGACATCCGCAACG
45	TGGCTCCGGAGGAGG	CGCCGCGCTCTCGCT	CGACATCCGCAACGA
	GGCTCCGGAGGAGGG	GCCGCGCTCTCGCTC	GACATCCGCAACGAC
	GCTCCGGAGGAGGGT	CCGCGCTCTCGCTCT	ACATCCGCAACGACT
	CTCCGGAGGAGGGTC	CGCGCTCTCGCTCTG	CATCCGCAACGACTA
	TCCGGAGGAGGGTCC	GCGCTCTCGCTCTGG	ATCCGCAACGACTAT
50	CCGGAGGAGGGTCCC	CGCTCTCGCTCTGGC	TCCGCAACGACTATC
	CGGAGGAGGGTCCCC	GCTCTCGCTCTGGCC	CCGCAACGACTATCA

CGCAACGACTATCAG	GGCTACCTCCACATC	CGCTTCCCCAAGCTC
GCAACGACTATCAGC	GCTACCTCCACATCC	GCTTCCCCAAGCTCA
CAACGACTATCAGCA	CTACCTCCACATCCT	CTTCCCCAAGCTCAC
AACGACTATCAGCAG	TACCTCCACATCCTG	TTCCCCAAGCTCACG
ACGACTATCAGCAGC	ACCTCCACATCCTGC	TCCCCAAGCTCACGG
CGACTATCAGCAGCT	CCTCCACATCCTGCT	CCCCAAGCTCACGGT
GACTATCAGCAGCTG	CTCCACATCCTGCTC	CCCAAGCTCACGGTC
ACTATCAGCAGCTGA	TCCACATCCTGCTCA	CCAAGCTCACGGTCA
CTATCAGCAGCTGAA	CCACATCCTGCTCAT	CAAGCTCACGGTCAT
TATCAGCAGCTGAAG	CACATCCTGCTCATC	AAGCTCACGGTCATT
ATCAGCAGCTGAAGC	ACATCCTGCTCATCT	AGCTCACGGTCATTA
TCAGCAGCTGAAGCG	CATCCTGCTCATCTC	GCTCACGGTCATTAC
CAGCAGCTGAAGCGC	ATCCTGCTCATCTCC	CTCACGGTCATTACC
AGCAGCTGAAGCGCC	TCCTGCTCATCTCCA	TCACGGTCATTACCG
GCAGCTGAAGCGCCT	CCTGCTCATCTCCAA	CACGGTCATTACCGA
CAGCTGAAGCGCCTG	CTGCTCATCTCCAAG	ACGGTCATTACCGAG
AGCTGAAGCGCCTGG	TGCTCATCTCCAAGG	CGGTCATTACCGAGT
GCTGAAGCGCCTGGA	GCTCATCTCCAAGGC	GGTCATTACCGAGTA
CTGAAGCGCCTGGAG	CTCATCTCCAAGGCC	GTCATTACCGAGTAC
TGAAGCGCCTGGAGA	TCATCTCCAAGGCCG	TCATTACCGAGTACT
GAAGCGCCTGGAGAA	CATCTCCAAGGCCGA	CATTACCGAGTACTT
AAGCGCCTGGAGAAC	ATCTCCAAGGCCGAG	ATTACCGAGTACTTG
AGCGCCTGGAGAACT	TCTCCAAGGCCGAGG	TTACCGAGTACTTGC
GCGCCTGGAGAACTG	CTCCAAGGCCGAGGA	TACCGAGTACTTGCT
CGCCTGGAGAACTGC	TCCAAGGCCGAGGAC	ACCGAGTACTTGCTG
GCCTGGAGAACTGCA	CCAAGGCCGAGGACT	CCGAGTACTTGCTGC
CCTGGAGAACTGCAC	CAAGGCCGAGGACTA	CGAGTACTTGCTGCT
CTGGAGAACTGCACG	AAGGCCGAGGACTAC	GAGTACTTGCTGCTG
TGGAGAACTGCACGG	AGGCCGAGGACTACC	AGTACTTGCTGCTGT
GGAGAACTGCACGGT	GGCCGAGGACTACCG	GTACTTGCTGCTGTT
GAGAACTGCACGGTG	GCCGAGGACTACCGC	TACTTGCTGCTGTTC
AGAACTGCACGGTGA	CCGAGGACTACCGCA	ACTTGCTGCTGTTCC
GAACTGCACGGTGAT	CGAGGACTACCGCAG	CTTGCTGCTGTTCCG
AACTGCACGGTGATC	GAGGACTACCGCAGC	TTGCTGCTGTTCCGA
ACTGCACGGTGATCG	AGGACTACCGCAGCT	TGCTGCTGTTCCGAG
CTGCACGGTGATCGA	GGACTACCGCAGCTA	GCTGCTGTTCCGAGT
TGCACGGTGATCGAG	GACTACCGCAGCTAC	CTGCTGTTCCGAGTG
GCACGGTGATCGAGG	ACTACCGCAGCTACC	TGCTGTTCCGAGTGG
CACGGTGATCGAGGG	CTACCGCAGCTACCG	GCTGTTCCGAGTGGC
ACGGTGATCGAGGGC	TACCGCAGCTACCGC	CTGTTCCGAGTGGCT
CGGTGATCGAGGGCT	ACCGCAGCTACCGCT	TGTTCCGAGTGGCTG
GGTGATCGAGGGCTA	CCGCAGCTACCGCTT	GTTCCGAGTGGCTGG
GTGATCGAGGGCTAC	CGCAGCTACCGCTTC	TTCCGAGTGGCTGGC
TGATCGAGGGCTACC	GCAGCTACCGCTTCC	TCCGAGTGGCTGGCC
GATCGAGGGCTACCT	CAGCTACCGCTTCCC	CCGAGTGGCTGGCCT
ATCGAGGGCTACCTC	AGCTACCGCTTCCCC	CGAGTGGCTGGCCTC
TCGAGGGCTACCTCC	GCTACCGCTTCCCCA	GAGTGGCTGGCCTCG
CGAGGGCTACCTCCA	CTACCGCTTCCCCAA	AGTGGCTGGCCTCGA
GAGGGCTACCTCCAC	TACCGCTTCCCCAAG	GTGGCTGGCCTCGAG
AGGGCTACCTCCACA	ACCGCTTCCCCAAGC	TGGCTGGCCTCGAGA
GGGCTACCTCCACAT	CCGCTTCCCCAAGCT	GGCTGGCCTCGAGAG

	GCTGGCCTCGAGAGC	GGCTGGAAACTCTTC	CTCAAGGATATTGGG
	CTGGCCTCGAGAGCC	GCTGGAAACTCTTCT	TCAAGGATATTGGGC
	TGGCCTCGAGAGCCT	CTGGAAACTCTTCTA	CAAGGATATTGGGCT
	GGCCTCGAGAGCCTC	TGGAAACTCTTCTAC	AAGGATATTGGGCTT
5	GCCTCGAGAGCCTCG	GGAAACTCTTCTACA	AGGATATTGGGCTTT
	CCTCGAGAGCCTCGG	GAAACTCTTCTACAA	GGATATTGGGCTTTA
	CTCGAGAGCCTCGGA	AAACTCTTCTACAAC	GATATTGGGCTTTAC
	TCGAGAGCCTCGGAG	AACTCTTCTACAACT	ATATTGGGCTTTACA
	CGAGAGCCTCGGAGA	ACTCTTCTACAACTA	TATTGGGCTTTACAA
10	GAGAGCCTCGGAGAC	CTCTTCTACAACTAC	ATTGGGCTTTACAAC
	AGAGCCTCGGAGACC	TCTTCTACAACTACG	TTGGGCTTTACAACC
	GAGCCTCGGAGACCT	CTTCTACAACTACGC	TGGGCTTTACAACCT
	AGCCTCGGAGACCTC	TTCTACAACTACGCC	GGGCTTTACAACCTG
	GCCTCGGAGACCTCT	TCTACAACTACGCCC	GGCTTTACAACCTGA
15	CCTCGGAGACCTCTT	CTACAACTACGCCCT	GCTTTACAACCTGAG
	CTCGGAGACCTCTTC	TACAACTACGCCCTG	CTTTACAACCTGAGG
	TCGGAGACCTCTTCC	ACAACTACGCCCTGG	TTTACAACCTGAGGA
	CGGAGACCTCTTCCC	CAACTACGCCCTGGT	TTACAACCTGAGGAA
	GGAGACCTCTTCCCC	AACTACGCCCTGGTC	TACAACCTGAGGAAC
20	GAGACCTCTTCCCCA	ACTACGCCCTGGTCA	ACAACCTGAGGAACA
	AGACCTCTTCCCCAA	CTACGCCCTGGTCAT	CAACCTGAGGAACAT
	GACCTCTTCCCCAAC	TACGCCCTGGTCATC	AACCTGAGGAACATT
	ACCTCTTCCCCAACC	ACGCCCTGGTCATCT	ACCTGAGGAACATTA
	CCTCTTCCCCAACCT	CGCCCTGGTCATCTT	CCTGAGGAACATTAC
25	CTCTTCCCCAACCTC	GCCCTGGTCATCTTC	CTGAGGAACATTACT
	TCTTCCCCAACCTCA	CCCTGGTCATCTTCG	TGAGGAACATTACTC
	CTTCCCCAACCTCAC	CCTGGTCATCTTCGA	GAGGAACATTACTCG
	TTCCCCAACCTCACG	CTGGTCATCTTCGAG	AGGAACATTACTCGG
	TCCCCAACCTCACGG	TGGTCATCTTCGAGA	GGAACATTACTCGGG
30	CCCCAACCTCACGGT	GGTCATCTTCGAGAT	GAACATTACTCGGGG
	CCCAACCTCACGGTC	GTCATCTTCGAGATG	AACATTACTCGGGGG
	CCAACCTCACGGTCA	TCATCTTCGAGATGA	ACATTACTCGGGGGG
	CAACCTCACGGTCAT	CATCTTCGAGATGAC	CATTACTCGGGGGGC
	AACCTCACGGTCATC	ATCTTCGAGATGACC	ATTACTCGGGGGGCC
35	ACCTCACGGTCATCC	TCTTCGAGATGACCA	TTACTCGGGGGGCCA
	CCTCACGGTCATCCG	CTTCGAGATGACCAA	TACTCGGGGGGCCAT
	CTCACGGTCATCCGC	TTCGAGATGACCAAT	ACTCGGGGGGCCATC
	TCACGGTCATCCGCG	TCGAGATGACCAATC	CTCGGGGGGCCATCA
	CACGGTCATCCGCGG	CGAGATGACCAATCT	TCGGGGGGCCATCAG
40	ACGGTCATCCGCGGC	GAGATGACCAATCTC	CGGGGGGCCATCAGG
	CGGTCATCCGCGGCT	AGATGACCAATCTCA	GGGGGGCCATCAGGA
	GGTCATCCGCGGCTG	GATGACCAATCTCAA	GGGGGCCATCAGGAT
	GTCATCCGCGGCTGG	ATGACCAATCTCAAG	GGGGCCATCAGGATT
	TCATCCGCGGCTGGA	TGACCAATCTCAAGG	GGGCCATCAGGATTG
45	CATCCGCGGCTGGAA	GACCAATCTCAAGGA	GGCCATCAGGATTGA
	ATCCGCGGCTGGAAA	ACCAATCTCAAGGAT	GCCATCAGGATTGAG
	TCCGCGGCTGGAAAC	CCAATCTCAAGGATA	CCATCAGGATTGAGA
	CCGCGGCTGGAAACT	CAATCTCAAGGATAT	CATCAGGATTGAGAA
	CGCGGCTGGAAACTC	AATCTCAAGGATATT	ATCAGGATTGAGAAA
50	GCGGCTGGAAACTCT	ATCTCAAGGATATTG	TCAGGATTGAGAAAA
	CGGCTGGAAACTCTT	TCTCAAGGATATTGG	CAGGATTGAGAAAAA

AGGATTGAGAAAAAT	CTGATCCTGGATGCG	AAGGAATGTGGGGAC
GGATTGAGAAAAATG	TGATCCTGGATGCGG	AGGAATGTGGGGACC
GATTGAGAAAAATGC	GATCCTGGATGCGGT	GGAATGTGGGGACCT
ATTGAGAAAAATGCT	ATCCTGGATGCGGTG	GAATGTGGGGACCTG
TTGAGAAAAATGCTG	TCCTGGATGCGGTGT	AATGTGGGGACCTGT
TGAGAAAAATGCTGA	CCTGGATGCGGTGTC	ATGTGGGGACCTGTG
GAGAAAAATGCTGAC	CTGGATGCGGTGTCC	TGTGGGGACCTGTGT
AGAAAAATGCTGACC	TGGATGCGGTGTCCA	GTGGGGACCTGTGTC
GAAAAATGCTGACCT	GGATGCGGTGTCCAA	TGGGGACCTGTGTCC
AAAAATGCTGACCTC	GATGCGGTGTCCAAT	GGGGACCTGTGTCCA
AAAATGCTGACCTCT	ATGCGGTGTCCAATA	GGGACCTGTGTCCAG
AAATGCTGACCTCTG	TGCGGTGTCCAATAA	GGACCTGTGTCCAGG
AATGCTGACCTCTGT	GCGGTGTCCAATAAC	GACCTGTGTCCAGGG
ATGCTGACCTCTGTT	CGGTGTCCAATAACT	ACCTGTGTCCAGGGA
TGCTGACCTCTGTTA	GGTGTCCAATAACTA	CCTGTGTCCAGGGAC
GCTGACCTCTGTTAC	GTGTCCAATAACTAC	CTGTGTCCAGGGACC
CTGACCTCTGTTACC	TGTCCAATAACTACA	TGTGTCCAGGGACCA
TGACCTCTGTTACCT	GTCCAATAACTACAT	GTGTCCAGGGACCAT
GACCTCTGTTACCTC	TCCAATAACTACATT	TGTCCAGGGACCATG
ACCTCTGTTACCTCT	CCAATAACTACATTG	GTCCAGGGACCATGG
CCTCTGTTACCTCTC	CAATAACTACATTGT	TCCAGGGACCATGGA
CTCTGTTACCTCTCC	AATAACTACATTGTG	CCAGGGACCATGGAG
TCTGTTACCTCTCCA	ATAACTACATTGTGG	CAGGGACCATGGAGG
CTGTTACCTCTCCAC	TAACTACATTGTGGG	AGGGACCATGGAGGA
TGTTACCTCTCCACT	AACTACATTGTGGGG	GGGACCATGGAGGAG
GTTACCTCTCCACTG	ACTACATTGTGGGGA	GGACCATGGAGGAGA
TTACCTCTCCACTGT	CTACATTGTGGGGAA	GACCATGGAGGAGAA
TACCTCTCCACTGTG	TACATTGTGGGGAAT	ACCATGGAGGAGAAG
ACCTCTCCACTGTGG	ACATTGTGGGGAATA	CCATGGAGGAGAAGC
CCTCTCCACTGTGGA	CATTGTGGGGAATAA	CATGGAGGAGAAGCC
CTCTCCACTGTGGAC	ATTGTGGGGAATAAG	ATGGAGGAGAAGCCG
TCTCCACTGTGGACT	TTGTGGGGAATAAGC	TGGAGGAGAAGCCGA
CTCCACTGTGGACTG	TGTGGGGAATAAGCC	GGAGGAGAAGCCGAT
TCCACTGTGGACTGG	GTGGGGAATAAGCCC	GAGGAGAAGCCGATG
CCACTGTGGACTGGT	TGGGGAATAAGCCCC	AGGAGAAGCCGATGT
CACTGTGGACTGGTC	GGGGAATAAGCCCCC	GGAGAAGCCGATGTG
ACTGTGGACTGGTCC	GGGAATAAGCCCCCA	GAGAAGCCGATGTGT
CTGTGGACTGGTCCC	GGAATAAGCCCCCAA	AGAAGCCGATGTGTG
TGTGGACTGGTCCCT	GAATAAGCCCCCAAA	GAAGCCGATGTGTGA
GTGGACTGGTCCCTG	AATAAGCCCCCAAAG	AAGCCGATGTGTGAG
TGGACTGGTCCCTGA	ATAAGCCCCCAAAGG	AGCCGATGTGTGAGA
GGACTGGTCCCTGAT	TAAGCCCCCAAAGGA	GCCGATGTGTGAGAA
GACTGGTCCCTGATC	AAGCCCCCAAAGGAA	CCGATGTGTGAGAAG
ACTGGTCCCTGATCC	AGCCCCCAAAGGAAT	CGATGTGTGAGAAGA
CTGGTCCCTGATCCT	GCCCCCAAAGGAATG	GATGTGTGAGAAGAC
TGGTCCCTGATCCTG	CCCCCAAAGGAATGT	ATGTGTGAGAAGACC
GGTCCCTGATCCTGG	CCCCAAAGGAATGTG	TGTGTGAGAAGACCA
GTCCCTGATCCTGGA	CCCAAAGGAATGTGG	GTGTGAGAAGACCAC
TCCCTGATCCTGGAT	CCAAAGGAATGTGGG	TGTGAGAAGACCACC
CCCTGATCCTGGATG	CAAAGGAATGTGGGG	GTGAGAAGACCACCA
CCTGATCCTGGATGC	AAAGGAATGTGGGGA	TGAGAAGACCACCAT

GAGAAGACCACCATC	CGCTGCCAGAAAATG	AACAATGAGTGCTGC
AGAAGACCACCATCA	GCTGCCAGAAAATGT	ACAATGAGTGCTGCC
GAAGACCACCATCAA	CTGCCAGAAAATGTG	CAATGAGTGCTGCCA
AAGACCACCATCAAC	TGCCAGAAAATGTGC	AATGAGTGCTGCCAC
AGACCACCATCAACA	GCCAGAAAATGTGCC	ATGAGTGCTGCCACC
GACCACCATCAACAA	CCAGAAAATGTGCCC	TGAGTGCTGCCACCC
ACCACCATCAACAAT	CAGAAAATGTGCCCA	GAGTGCTGCCACCCC
CCACCATCAACAATG	AGAAAATGTGCCCAA	AGTGCTGCCACCCCG
CACCATCAACAATGA	GAAAATGTGCCCAAG	GTGCTGCCACCCCGA
ACCATCAACAATGAG	AAAATGTGCCCAAGC	TGCTGCCACCCCGAG
CCATCAACAATGAGT	AAATGTGCCCAAGCA	GCTGCCACCCCGAGT
CATCAACAATGAGTA	AATGTGCCCAAGCAC	CTGCCACCCCGAGTG
ATCAACAATGAGTAC	ATGTGCCCAAGCACG	TGCCACCCCGAGTGC
TCAACAATGAGTACA	TGTGCCCAAGCACGT	GCCACCCCGAGTGCC
CAACAATGAGTACAA	GTGCCCAAGCACGTG	CCACCCCGAGTGCCT
AACAATGAGTACAAC	TGCCCAAGCACGTGT	CACCCCGAGTGCCTG
ACAATGAGTACAACT	GCCCAAGCACGTGTG	ACCCCGAGTGCCTGG
CAATGAGTACAACTA	CCCAAGCACGTGTGG	CCCCGAGTGCCTGGG
AATGAGTACAACTAC	CCAAGCACGTGTGGG	CCCGAGTGCCTGGGC
ATGAGTACAACTACC	CAAGCACGTGTGGGA	CCGAGTGCCTGGGCA
TGAGTACAACTACCG	AAGCACGTGTGGGAA	CGAGTGCCTGGGCAG
GAGTACAACTACCGC	AGCACGTGTGGGAAG	GAGTGCCTGGGCAGC
AGTACAACTACCGCT	GCACGTGTGGGAAGC	AGTGCCTGGGCAGCT
GTACAACTACCGCTG	CACGTGTGGGAAGCG	GTGCCTGGGCAGCTG
TACAACTACCGCTGC	ACGTGTGGGAAGCGG	TGCCTGGGCAGCTGC
ACAACTACCGCTGCT	CGTGTGGGAAGCGGG	GCCTGGGCAGCTGCA
CAACTACCGCTGCTG	GTGTGGGAAGCGGGC	CCTGGGCAGCTGCAG
AACTACCGCTGCTGG	TGTGGGAAGCGGGCG	CTGGGCAGCTGCAGC
ACTACCGCTGCTGGA	GTGGGAAGCGGGCGT	TGGGCAGCTGCAGCG
CTACCGCTGCTGGAC	TGGGAAGCGGGCGTG	GGGCAGCTGCAGCGC
TACCGCTGCTGGACC	GGGAAGCGGGCGTGC	GGCAGCTGCAGCGCG
ACCGCTGCTGGACCA	GGAAGCGGGCGTGCA	GCAGCTGCAGCGCGC
CCGCTGCTGGACCAC	GAAGCGGGCGTGCAC	CAGCTGCAGCGCGCC
CGCTGCTGGACCACA	AAGCGGGCGTGCACC	AGCTGCAGCGCGCCT
GCTGCTGGACCACAA	AGCGGGCGTGCACCG	GCTGCAGCGCGCCTG
CTGCTGGACCACAAA	GCGGGCGTGCACCGA	CTGCAGCGCGCCTGA
TGCTGGACCACAAAC	CGGGCGTGCACCGAG	TGCAGCGCGCCTGAC
GCTGGACCACAAACC	GGGCGTGCACCGAGA	GCAGCGCGCCTGACA
CTGGACCACAAACCG	GGCGTGCACCGAGAA	CAGCGCGCCTGACAA
TGGACCACAAACCGC	GCGTGCACCGAGAAC	AGCGCGCCTGACAAC
GGACCACAAACCGCT	CGTGCACCGAGAACA	GCGCGCCTGACAACG
GACCACAAACCGCTG	GTGCACCGAGAACAA	CGCGCCTGACAACGA
ACCACAAACCGCTGC	TGCACCGAGAACAAT	GCGCCTGACAACGAC
CCACAAACCGCTGCC	GCACCGAGAACAATG	CGCCTGACAACGACA
CACAAACCGCTGCCA	CACCGAGAACAATGA	GCCTGACAACGACAC
ACAAACCGCTGCCAG	ACCGAGAACAATGAG	CCTGACAACGACACG
CAAACCGCTGCCAGA	CCGAGAACAATGAGT	CTGACAACGACACGG
AAACCGCTGCCAGAA	CGAGAACAATGAGTG	TGACAACGACACGGC
AACCGCTGCCAGAAA	GAGAACAATGAGTGC	GACAACGACACGGCC
ACCGCTGCCAGAAAA	AGAACAATGAGTGCT	ACAACGACACGGCCT
CCGCTGCCAGAAAAT	GAACAATGAGTGCTG	CAACGACACGGCCTG

	AACGACACGGCCTGT	GTGCCTGCCTGCCCG	GACCGTGACTTCTGC
	ACGACACGGCCTGTG	TGCCTGCCTGCCCGC	ACCGTGACTTCTGCG
	CGACACGGCCTGTGT	GCCTGCCTGCCCGCC	CCGTGACTTCTGCGC
	GACACGGCCTGTGTA	CCTGCCTGCCCGCCC	CGTGACTTCTGCGCC
5	ACACGGCCTGTGTAG	CTGCCTGCCCGCCCA	GTGACTTCTGCGCCA
	CACGGCCTGTGTAGC	TGCCTGCCCGCCCAA	TGACTTCTGCGCCAA
	ACGGCCTGTGTAGCT	GCCTGCCCGCCCAAC	GACTTCTGCGCCAAC
	CGGCCTGTGTAGCTT	CCTGCCCGCCCAACA	ACTTCTGCGCCAACA
	GGCCTGTGTAGCTTG	CTGCCCGCCCAACAC	CTTCTGCGCCAACAT
10	GCCTGTGTAGCTTGC	TGCCCGCCCAACACC	TTCTGCGCCAACATC
	CCTGTGTAGCTTGCC	GCCCGCCCAACACCT	TCTGCGCCAACATCC
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15	ATTACAAAAAAACAC	TCTAGGGACATGAAA	ATTTTAACGCTGCCT
	TTACAAAAAAACACG	CTAGGGACATGAAAT	TTTTAACGCTGCCTA
	TACAAAAAAACACGT	TAGGGACATGAAATT	TTTAACGCTGCCTAA
	ACAAAAAAACACGTG	AGGGACATGAAATTT	TTAACGCTGCCTAAT
	CAAAAAAACACGTGG	GGGACATGAAATTTA	TAACGCTGCCTAATT
20	AAAAAAACACGTGGA	GGACATGAAATTTAC	AACGCTGCCTAATTT
	AAAAAACACGTGGAG	GACATGAAATTTACA	ACGCTGCCTAATTTT
	AAAAACACGTGGAGA	ACATGAAATTTACAA	CGCTGCCTAATTTTG
	AAAACACGTGGAGAT	CATGAAATTTACAAA	GCTGCCTAATTTTGC
	AAACACGTGGAGATG	ATGAAATTTACAAAG	CTGCCTAATTTTGCC
25	AACACGTGGAGATGG	TGAAATTTACAAAGG	TGCCTAATTTTGCCA
	ACACGTGGAGATGGA	GAAATTTACAAAGGG	GCCTAATTTTGCCAA
	CACGTGGAGATGGAA	AAATTTACAAAGGGC	CCTAATTTTGCCAAA
	ACGTGGAGATGGAAA	AATTTACAAAGGGCC	CTAATTTTGCCAAAA
	CGTGGAGATGGAAAT	ATTTACAAAGGGCCA	TAATTTTGCCAAAAT
30	GTGGAGATGGAAATT	TTTACAAAGGGCCAT	AATTTTGCCAAAATC
	TGGAGATGGAAATTT	TTACAAAGGGCCATC	ATTTTGCCAAAATCC
	GGAGATGGAAATTTT	TACAAAGGGCCATCG	TTTTGCCAAAATCCT
	GAGATGGAAATTTTT	ACAAAGGGCCATCGT	TTTGCCAAAATCCTG
	AGATGGAAATTTTTA	CAAAGGGCCATCGTT	TTGCCAAAATCCTGA
35	GATGGAAATTTTTAC	AAAGGGCCATCGTTC	TGCCAAAATCCTGAA
	ATGGAAATTTTTACC	AAGGGCCATCGTTCA	GCCAAAATCCTGAAC
	TGGAAATTTTTACCT	AGGGCCATCGTTCAT	CCAAAATCCTGAACT
	GGAAATTTTTACCTT	GGGCCATCGTTCATC	CAAAATCCTGAACTT
	GAAATTTTTACCTTT	GGCCATCGTTCATCC	AAAATCCTGAACTTT
40	AAATTTTTACCTTTA	GCCATCGTTCATCCA	AAATCCTGAACTTTC
	AATTTTTACCTTTAT	CCATCGTTCATCCAA	AATCCTGAACTTTCT
	ATTTTTACCTTTATC	CATCGTTCATCCAAG	ATCCTGAACTTTCTC
	TTTTTACCTTTATCT	ATCGTTCATCCAAGG	TCCTGAACTTTCTCC
	TTTTACCTTTATCTT	TCGTTCATCCAAGGC	CCTGAACTTTCTCCC
45	TTTACCTTTATCTTT	CGTTCATCCAAGGCT	CTGAACTTTCTCCCT
	TTACCTTTATCTTTC	GTTCATCCAAGGCTG	TGAACTTTCTCCCTC
	TACCTTTATCTTTCA	TTCATCCAAGGCTGT	GAACTTTCTCCCTCA
	ACCTTTATCTTTCAC	TCATCCAAGGCTGTT	AACTTTCTCCCTCAT
	CCTTTATCTTTCACC	CATCCAAGGCTGTTA	ACTTTCTCCCTCATC
50	CTTTATCTTTCACCT	ATCCAAGGCTGTTAC	CTTTCTCCCTCATCG
	TTTATCTTTCACCTT	TCCAAGGCTGTTACC	TTTCTCCCTCATCGG

	TTCTCCCTCATCGGC	GCATGGCAGCTGGTT	CCATCCGACTGCCCC
	TCTCCCTCATCGGCC	CATGGCAGCTGGTTG	CATCCGACTGCCCCT
	CTCCCTCATCGGCCC	ATGGCAGCTGGTTGC	ATCCGACTGCCCCTG
	TCCCTCATCGGCCCG	TGGCAGCTGGTTGCT	TCCGACTGCCCCTGC
5	CCCTCATCGGCCCGG	GGCAGCTGGTTGCTC	CCGACTGCCCCTGCT
	CCTCATCGGCCCGGC	GCAGCTGGTTGCTCC	CGACTGCCCCTGCTG
	CTCATCGGCCCGGCG	CAGCTGGTTGCTCCA	GACTGCCCCTGCTGT
	TCATCGGCCCGGCGC	AGCTGGTTGCTCCAT	ACTGCCCCTGCTGTG
	CATCGGCCCGGCGCT	GCTGGTTGCTCCATT	CTGCCCCTGCTGTGC
10	ATCGGCCCGGCGCTG	CTGGTTGCTCCATTT	TGCCCCTGCTGTGCT
	TCGGCCCGGCGCTGA	TGGTTGCTCCATTTG	GCCCCTGCTGTGCTG
	CGGCCCGGCGCTGAT	GGTTGCTCCATTTGA	CCCCTGCTGTGCTGC
	GGCCCGGCGCTGATT	GTTGCTCCATTTGAG	CCCTGCTGTGCTGCT
	GCCCGGCGCTGATTC	TTGCTCCATTTGAGA	CCTGCTGTGCTGCTC
15	CCCGGCGCTGATTCC	TGCTCCATTTGAGAG	CTGCTGTGCTGCTCA
	CCGGCGCTGATTCCT	GCTCCATTTGAGAGA	TGCTGTGCTGCTCAA
	CGGCGCTGATTCCTC	CTCCATTTGAGAGAC	GCTGTGCTGCTCAAG
	GGCGCTGATTCCTCG	TCCATTTGAGAGACA	CTGTGCTGCTCAAGG
	GCGCTGATTCCTCGT	CCATTTGAGAGACAC	TGTGCTGCTCAAGGC
20	CGCTGATTCCTCGTG	CATTTGAGAGACACG	GTGCTGCTCAAGGCC
	GCTGATTCCTCGTGT	ATTTGAGAGACACGC	TGCTGCTCAAGGCCA
	CTGATTCCTCGTGTC	TTTGAGAGACACGCT	GCTGCTCAAGGCCAC
	TGATTCCTCGTGTCC	TTGAGAGACACGCTG	CTGCTCAAGGCCACA
	GATTCCTCGTGTCCG	TGAGAGACACGCTGG	TGCTCAAGGCCACAG
25	ATTCCTCGTGTCCGG	GAGAGACACGCTGGC	GCTCAAGGCCACAGG
	TTCCTCGTGTCCGGA	AGAGACACGCTGGCG	CTCAAGGCCACAGGC
	TCCTCGTGTCCGGAG	GAGACACGCTGGCGA	TCAAGGCCACAGGCA
	CCTCGTGTCCGGAGG	AGACACGCTGGCGAC	CAAGGCCACAGGCAC
	CTCGTGTCCGGAGGC	GACACGCTGGCGACA	AAGGCCACAGGCACA
30	TCGTGTCCGGAGGCA	ACACGCTGGCGACAC	AGGCCACAGGCACAC
	CGTGTCCGGAGGCAT	CACGCTGGCGACACA	GGCCACAGGCACACA
	GTGTCCGGAGGCATG	ACGCTGGCGACACAC	GCCACAGGCACACAG
	TGTCCGGAGGCATGG	CGCTGGCGACACACT	CCACAGGCACACAGG
	GTCCGGAGGCATGGG	GCTGGCGACACACTC	CACAGGCACACAGGT
35	TCCGGAGGCATGGGT	CTGGCGACACACTCC	ACAGGCACACAGGTC
	CCGGAGGCATGGGTG	TGGCGACACACTCCG	CAGGCACACAGGTCT
	CGGAGGCATGGGTGA	GGCGACACACTCCGT	AGGCACACAGGTCTC
	GGAGGCATGGGTGAG	GCGACACACTCCGTC	GGCACACAGGTCTCA
	GAGGCATGGGTGAGC	CGACACACTCCGTCC	GCACACAGGTCTCAT
40	AGGCATGGGTGAGCA	GACACACTCCGTCCA	CACACAGGTCTCATT
	GGCATGGGTGAGCAT	ACACACTCCGTCCAT	ACACAGGTCTCATTG
	GCATGGGTGAGCATG	CACACTCCGTCCATC	CACAGGTCTCATTGC
	CATGGGTGAGCATGG	ACACTCCGTCCATCC	ACAGGTCTCATTGCT
	ATGGGTGAGCATGGC	CACTCCGTCCATCCG	CAGGTCTCATTGCTT
45	TGGGTGAGCATGGCA	ACTCCGTCCATCCGA	AGGTCTCATTGCTTC
	GGGTGAGCATGGCAG	CTCCGTCCATCCGAC	GGTCTCATTGCTTCT
	GGTGAGCATGGCAGC	TCCGTCCATCCGACT	GTCTCATTGCTTCTG
	GTGAGCATGGCAGCT	CCGTCCATCCGACTG	TCTCATTGCTTCTGA
	TGAGCATGGCAGCTG	CGTCCATCCGACTGC	CTCATTGCTTCTGAC
50	GAGCATGGCAGCTGG	GTCCATCCGACTGCC	TCATTGCTTCTGACT
	AGCATGGCAGCTGGT	TCCATCCGACTGCCC	CATTGCTTCTGACTA

ATTGCTTCTGACTAG	CTCTCAGTGAAGGTG
TTGCTTCTGACTAGA	TCTCAGTGAAGGTGG
TGCTTCTGACTAGAT	CTCAGTGAAGGTGGG
GCTTCTGACTAGATT	TCAGTGAAGGTGGGG
CTTCTGACTAGATTA	CAGTGAAGGTGGGGA
TTCTGACTAGATTAT	AGTGAAGGTGGGGAG
TCTGACTAGATTATT	GTGAAGGTGGGGAGA
CTGACTAGATTATTA	TGAAGGTGGGGAGAA
TGACTAGATTATTAT	GAAGGTGGGGAGAAG
GACTAGATTATTATT	AAGGTGGGGAGAAGC
ACTAGATTATTATTT	AGGTGGGGAGAAGCT
CTAGATTATTATTTG	GGTGGGGAGAAGCTG
TAGATTATTATTTGG	GTGGGGAGAAGCTGA
AGATTATTATTTGGG	TGGGGAGAAGCTGAA
GATTATTATTTGGGG	GGGGAGAAGCTGAAC
ATTATTATTTGGGGG	GGGAGAAGCTGAACC
TTATTATTTGGGGGA	GGAGAAGCTGAACCG
TATTATTTGGGGGAA	GAGAAGCTGAACCGG
ATTATTTGGGGGAAC	AGAAGCTGAACCGGC
TTATTTGGGGGAACT	
TATTTGGGGGAACTG	
ATTTGGGGGAACTGG	
TTTGGGGGAACTGGA	
TTGGGGGAACTGGAC	
TGGGGGAACTGGACA	
GGGGGAACTGGACAC	
GGGGAACTGGACACA	
GGGAACTGGACACAA	
GGAACTGGACACAAT	
GAACTGGACACAATA	
AACTGGACACAATAG	
ACTGGACACAATAGG	
CTGGACACAATAGGT	
TGGACACAATAGGTC	
GGACACAATAGGTCT	
GACACAATAGGTCTT	
ACACAATAGGTCTTT	
CACAATAGGTCTTTC	
ACAATAGGTCTTTCT	
CAATAGGTCTTTCTC	
AATAGGTCTTTCTCT	
ATAGGTCTTTCTCTC	
TAGGTCTTTCTCTCA	
AGGTCTTTCTCTCAG	
GGTCTTTCTCTCAGT	
GTCTTTCTCTCAGTG	
TCTTTCTCTCAGTGA	
CTTTCTCTCAGTGAA	
TTTCTCTCAGTGAAG	
TTCTCTCAGTGAAGG	
TCTCTCAGTGAAGGT	

EXAMPLE 9

Sub-confluent HaCaT cells were treated as described above with phosphorothioate oligonucleotides IGFR.AS (antisense: 5′-ATCTCTCCGCTTCCTTTC-3′; (<400>10); ref 13) and IGFR.S (sense control: 5′-GAAAGGAAGCGGAGAGAT-3′; (<400>11); ref 13) IGF-I binding to the cell monolayers was then measured as ^{125}I-IGF-I.

EXAMPLE 10

The results of this experiment are shown in Figures 7 and 8.

HaCaT cells were initially plated in DMEM with 10% v/v serum, then AS oligo experiments were performed in complete "Keratinocyte-SFM" (Gibco) to exclude the influence of exogenous IGFBPs. Oligos were synthesised as phosphorothioate (nuclease-resistant) derivatives (Bresatec, South Australia) and were as follows: antisense: AS2, 5′-GCGCCCGCTGCATGACGCCTGCAAC-3′ (IGFBP-3 start codon); controls: AS2NS, 5′-CGGAGATGCCGCATGCCAGCGCAGG-3′; AS4, 5′-AGGCGGCTGACGGCACTA-3′; AS4NS, 5′-GACAGCGTCGGAGCGATC-3′; IGFRAS, 5′-ATCTCTCCGCTTCCTTTC-3′; IGFRS, 5′-GAAAGGAAGCGGAGAGAT-3′. Oligos to IGFBP-3 were based on the published sequence of Spratt *et al* [12]. AS oligos were added to HaCaT monolayers in 0.5ml medium in 24-well plates at the concentrations and addition frequencies indicated. IGFBP-3 measured in cell-conditioned medium using a dot-blot assay, adapted from the Western ligand blot method of Hossenlopp *et al* [11], in which 100μl of conditioned medium was applied to nitrocellulose filters with a vacuum dot-blot apparatus. After drying the membranes at 37°C, relative amounts of IGFBP are determined by ^{125}I-IGF-I-binding, autoradiography and computerised imaging densitometry. Triplicate wells (except in Figure 7, where duplicate wells were measured as shown) were analysed and corrected for changes in cell number per well. Relative cell number per well was determined using an amido black dye method, developed specifically for cultured monolayers of HaCaT cells [14]. Cell numbers differed by less than 10% after treatment. For oligos to the IGF receptor, receptor quantitation in

intact HaCaT monolayers was by overnight incubation with ^{125}I-IGF-I (30,000cpm/well) at 4°C.

EXAMPLE 11

Experiments involving ribozymes are generally conducted as described in Internaitonal Patent Application No. WO 89/05852 and in Haselhoff and Gerlach [8]. Ribozymes are constructed with a hybridising region which is complementary in nucleotide sequence to at least part of a target RNA which, in this case, encodes IGFBP-2. Activity of ribozymes is measurable on, for example, Northern blots or using animal models such as in the nude mouse model (15; 16) or the "flaky skin" mouse model (17; 18).

EXAMPLE 12

The methods described in Example 11 are used for the screening of ribozymes which inhibit IGFBP-3 production. The activity of the ribozymes is determined as in Example 11.

EXAMPLE 13

The methods described in Example 11 are used for the screening of ribozymes which inhibit IGF-1 production. The activity of the ribozymes is determined as in Example 11.

EXAMPLE 14

The methods described in Example 11 are used for the screening of ribozymes which inhibit IGF-1 production. The activity of the ribozymes is determined as in Example 11.

EXAMPLE 15

Twenty-one antisense oligonucleotides targeted to mRNA sequences enducing the IGF-1 receptor, and four random oligonucleotides were synthesized. The antisense oligonucleotides are C5-propynyl-dU, dC 15mer phosphorothioate oligodeoxyribonucleotides. In these oligonucleotides, a phosphorothioate backbone replaces the phosphodiester backbone of naturally occurring DNA. The positions of the 21 sequence specific antisense oligonucleotides relative to the IGF-1 receptor mRNA structure are shown in Figure 9.

EXAMPLE 16

Experiments were performed to determine the uptake of the antisense oligonucleotides of Example 15 into keratinocytes. Cells of the differentiated human keratinocyte cell line, HaCaT, were incubated for 24 hours in Dulbecco's Modified Eagle Medium (DMEM) supplemented with 10% (w/v) fetal calf serum (FCS) containing fluorescently labelled oligonucleotide (R451, a randomized sequence oligonucleotide, 30nM) and cytofectin GSV (2μg/ml, Glen Research, 44901 Falcon Place, Sterling, VA 20166, Cat. No. 70-3815-78). Cells were then transferred to oligonucleotide-free medium and fluorescence microcopy and phase contrast images of the cells were obtained. Figure 10 shows fluorescence microscopy (Panel A) and phase contrast (Panel B) images of uptake of fluorescently labelled oligonucleotide in the majority of cells in a HaCaT monolayer. The degree of uptake obtained with the cationic lipid cytofectin was far greater than the uptake obtained with the next best lipid tried, Tfx-50.

A further experiment was performed to assess the uptake and toxicity associated with the use of cytofectin GSV over five days. Confluent HaCaT keratinocytes were incubated in DMEM containing fluorescently labelled oligonucleotide R451 (30nM or 100 nM) plus cytofectin GSV (2μg/ml or 5μg/ml) over 120 hours, viewed by fluorescence microscopy, tryptan blue stained, and counted. The graphs in Figure 11 depict uptake (Panel A) and toxicity (Panel B). The proportion of cells containing oligonucleotide remained high over the 120 hour period. The combination of 30 nM oligonucleotide and 2μg/ml GSV provided optimal uptake and minimal toxicity.

EXAMPLE 17

The twenty-one oligonucleotides of Example 15 were then screened for their ability to inhibit IGF-I receptor mRNA levels in HaCaT cells, in accordance with the teachings herein. HaCaT cells were grown to 90% confluence in DMEM supplemented with 10% (v/v) FCS. Antisense oligonucleotides (30nM) were completed with cytofectin GSV (2μg/ml) and added tot he cells in the presence of serum. HaCaT keratinocytes were treated with the oligonucleotide/GSV complexes or randomized sequence oligonucleotides (R451, R766),

liposome alone (GSV), or were left untreated (UT). Duplicate treatments were performed. Repeat additions of the oligonucleotides/GSV complex were performed at 24, 48 and 76 hours following the first addition. Total RNA was isolated as per the RNAzolB protocol (Biotecx Laboratories, Inc. 6023 South Loop East, Houston, TX 77033) 96 hours following the first addition.

IGF-I receptor mRNA and glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA levels were simultaneously determined by a ribonuclease (RNase) protection assay. The RNase Protection Assay kit, *in vitro* transcription kit, and IGF-I receptor and GAPDH DNA templates were obtained from Ambion, Inc. (2130 Woodward St., Houston, TX 78744). The amount of IGF-I receptor mRNA in any given sample was expressed as the amount of IGF-I receptor mRNA relative to the amount of GAPDH mRNA. Each oligonucleotide was tested in at least two separate experiments.

Figure 12 depicts representative results of the screening process. Panel A shows an electrophoretic analysis of IGF-I receptor and GAPDH mRNA fragments after RNase protection. Molecular weight markers are shown on the right hand side. The full-length probe is shown on the left hand side; G-probe indicates the IGF-I receptor probe. GAPDH protected fragments (G) are seen at 316 bases and IGF-I protected fragments (I) are seen at 276 bases. Exhibit E, Panel B provides a graph indicating the relative level of IGF-I receptor mRNA following each treatment.

The results obtaining from the above screening assays are summarized in Figure 13. The graph depicts the relative level of IGF-I receptor mRNA after treatment with oligonucleotides complementary to the human IGF-I receptor mRNA (26-86), four randomized sequence oligonucleotides (R1, R4, R7, R9), liposome alone (GSV), or no treatment (UT). Asterisks indicate a significant different in relative IGF-I receptor mRNA as compared to GSV treated cells ($n=4\text{-}10$, $p<0.05$).

As demonstrated in Figure 13, treatment with eighteen of the twenty-one oligonucleotides resulted in a significant different in levels of IGF-I receptor mRNA relative to GSV treated cells. Three of the antisense oligonucleotides tested in the screening assay reduce IGF-I receptor mRNA to less than 35% of GSV-treated cells. These antisense oligonucleotides have the following sequences, presented in the 5′ to 3′ direction:

#27 UCCGGAGCCAGACUU
#64 CACAGUUGCUGCAAG
#78 UCUCCGCUUCCUUUC

As further demonstrated in Figure 13, six of the antisense oligonucleotides tested in the screening assay reduce IGF-I receptor mRNA to between 35 and 50% of GSV-treated cells. These antisense oligonucleotides have the following sequences, presented in the 5′ to 3′ direction:

#28 AGCCCCCACAGCGAG
#32 GCCUUGGAGAUGAGC
#40 UAACAGAGGUCAGCA
#42 GGAUCAGGGACCAGU
#46 CGGCAAGCUACACAG
#50 GGCAGGCAGGCACAC

EXAMPLE 19

Another experiment was performed demonstrating that antisense oligonucleotides targeted to genetic sequences encoding the IGF0I receptor and that reduce IGF-I receptor mRNA levels also inhibit the IGF-I receptor level on the surface of the treated cultured keratinocytes. HaCaT cells were grown to confluence in 24-well plates in DMEM containing 10% (v/v) FCS. Oligodeoxynucleotide and cytofectin GSV were mixed together in serum-free DMEM, and incubated at room temperature for 10 minutes before being diluted ten-fold in medium and placed on the cells. Cells were incubated for 72 hours with 30nM random sequence or

antisense oligonucleotide and 2μm/ml GSV, or with GSV alone in DMEM containing 10% (v/v) FCS with solutions replaced every 24 hours. This was followed by incubation with oligonucleotide/GSV in serum-free DMEM for 48 hours. All incubations were performed at 37°C. Cells were washed twice with 1ml cold PBS. Serum-free DMEM containing 10^{-10}M ^{125}I-IGF-I was added with or without the IGF-I analogue, des (1-3) IGF-I, at 10^{-11}M to 10^{-7}M. Cells were incubated at 4°C for 17 hours with gentle shaking, then washed three times with 1ml cold PBS and lysed in 250μl 0.5M NaOH/0.1% (v/v) Triton X-100 at room temperature for 4 hours. Specific binding of the solubilised cell extract was measured using a gamma counter. As shown in Figure 14, treatment of HaCaT keratinocytes with oligonucleotide reduced cell surface IGF-I receptor levels to 30% of levels in untreated keratinocytes or in keratinocytes treated with liposome alone or a random oligonucleotide, R766. As shown in Figure 15, treatment with oligonucleotide #27 also significantly reduced cell surface IGF-I receptor levels relative to untreated keratinocytes or treatment with liposome alone or random nucleotide R451. As demonstrated in Example 17, oligonucleotides #64 and #27 reduce IGF-I receptor mRNA levels in cultured keratinocytes to less than 35% of GSV-treated cells. Accordingly, the ability of an oligonucleotide to reduce IGF-I receptor mRNA levels in correlated with its ability to reduce cell surface IGF-I receptor levels.

The forgoing Examples demonstrate that antisense oligonucleotides targeted to the IGF-I receptor can be delivered to human keratinocytes *in vitro*, can inhibit IGF-I receptor mRNA levels in human keratinocytes *in vitro*, and that inhibition of mRNA levels is correlated with reduction of cell surface IGF-I receptor levels.

EXAMPLE 19

Further experiments demonstrated the efficacy of antisense oligonucleotides targeted tot he IGF-I receptor in an *in vivo* model of psoriasis. An animal model of psoriasis is the human psoriatic skin xenograft model. The skin used in this model contains the true disease state. In this model, reduction in epidermal thickness of psoriatic grafts in response to treatment is positively correlated with efficacy of treatment. Both normal and psoriatic human skin were

grated into a thymic (nude) mice in accordance with a thymic (nude) mice in accordance with the methods of Baker *et al* (1992) *Brit. J. Dermatol.* 126:105 and Nanney *et al* (1992) *J. Invest. Dermatol,* 92:296. Successful grafting was achieved, as demonstrated in Figure 16, which shows hemotoxylin and eosin (H&E) stained sections of a 49-day old psoriatic human skin graft (Panel B) compared to the histology of the skin graft prior to grafting (Panel A). The histological features of psoriasis present in the pregraft section (e.g., parakeratosis, acanthosis and pronounced rete ridges) are present in the grafts more than seven weeks post grafting.

Using the model, oligonucleotide uptake was measured in epidermal keratinocytes *in vivo* after intradermal injection. Fluorescently labelled oligonucleotide (R451, 50μl, 10μM injection) was intradermally injected into psoriatic and normal skin grafts on a thymic mice. Live confocal microscopy and fluorescence microscopy of fixed sections was then employed. Using both techniques, oligonucleotide was found to localize in the nucleus of over 90% of basal keratinocytes. Figure 17 shows the nuclear localization of oligonucleotide in psoriatic skin cells using conventional fluorescence microscopy of a graft that was removed and sectioned after 24 hours.

After establishing oligonucleotide uptake in the *in vivo* model, a small number of pilots experiments were performed to determine a schedule for treatment of grated mice with antisense oligonucleotides targeted to genetic sequences encoding the IGF-I receptor. The treatment schedule was finalized as follows:

Graft Number	Treatment	Volume of Injection	ODN Concentration	Duration of Treatment
1-3	Vehicle (PBS)	50μl	-	20 days
4-6	RandomODN#R451	50μl	10μM	20 days
7-9	ODN#27	50μl	10μM	20 days
10-12	ODN#74	50μl	10μM	20 days
13-15	ODN#50	50μl	10μM	20 days

As determined above, oligonucleotide #27 (ODN #27) reduced IGF-I receptor mRNA *in vitro* to less than 35% of GSV-treated cells. Oligonucleotide #50 (ODN#50) reduced IGF-I receptor mRNA *in vitro* to between 35 and 50% of GSV-treated cells. Oligonucleotide #74 (ODN #74) was not inhibitory to IGF-I receptor mRNA *in vitro*. In the *in vivo* model, each mouse received two grafts. Random oligonucleotide or vehicle was injected intradermally in one graft and acted as a control. The second graft was injected with the targeted oligonucleotide. Each graft received an injection every second day for the duration of the treatment.

Histology of representative grafts from each treatment type are shown in Figures 18(a)-(d) and 19(a) - (d). Each sheet shows three images of H&E stained sections: the pregraft histology, the control treated graft, and the targeted oligonucleotide treated graft. Figures 18(a)-(d) are shown at 100x magnification; figures 19(a)-(d) are shown at 400x magnification. The total cross sectional area of epidermis of each graft was assessed using MCID analysis software. The pooled results from all of the treated grafts are shown in Figure 20.

As shown in Figures 18(a)-(d) and 19(a)-(d), the vehicle-treated (control) grafts were marginally thinner than thepregraft sections. The degree of regression in these experiments (ie., less than 10%) is not significant. A similar amount of marginal thinning

of epidermis compared to pregraft also occurred in pilot experiments in which psoriatic grafts were not injected, and thsu it is unlikely that the vehicle itself has any effect. Histological features of psoriasis present in skin samples prior to grafting (clubbing of rete ridges, parakeratosis, acanthosis) were present in these grafts.

The random oliognucleotide treated grafts varied in epidermal thickness after 20 days of treatment. Grafts were either a similar thickness to the pregraft histology, or marginally thinner. Random oligonucleotide treated grafts were in each case significantly thicker than their targeted oligonucleotide treated pairs.

As shown in Figure 20, the targeted oligonucleotide treated grafts were significantly thinner than the pregraft sections and showed less parakeratosis and clubbing of rete ridges. Antisense oligonucleotides which were effective at reducing IGF-I receptor mRNA levels *in vitro* (#27 and #50) produced greatere epidermal thinning than an oligonucleotide which was not inhibitory to IGF-I receptor mRNA *in vitro* (#74). Accordingly, there is a direct correlation between the ability of an oligonucleotide targeted to the IGF-I receptor to inhibit IGF-I receptor mRNA levels *in vitro* and the efficacy of the oligonucleotide as an anti-psoriasis agent in an *in vivo* model.

EXAMPLE 20

Another experiment demonstrated that treatment of psoriatic grafts with an oligonucleotide targeted to a genetic sequence encoding the IGF-I receptor results in inhibition of proliferation. Pregrafts from psoriatic patients, control grafts treated with R4541, and grafts treated with oligonucleotide #27 were obtained as described in Example 19. An antibody to the cell cycle-specific nuclear antigen Ki67 was used to immunohistochemically detect actively dividing cells and tereby assess proliferation. The αKi67 antibody (DAKO, Glostrup, Denmark) recognizes the Ki67 antigen transiently expressed in nuclei of proliferating cells during late G_1, S, M and G_2 phases of the cycle and thsu provides a marker for proliferation. Pregraft and graft sections were immunohistochemically processed by standard methods using αKi67 (according to the

manufacturer's instructions), peroxidase-conjugated anti-rabbit second stage antibody, and a chromogenic peroxidase substrate.

The results of this experiment are presented in Figure 21 as immunohistochemical sections at 100x magnification. The top panel of Figure 21 depicts a pregraft section obtained from a psoriatic patient. The epidermis is thicker than normal and nucleic are evident in the stratum corneum. Ki67 positive cells, appearing as brown dots, are evidence in the basal and suprabasal layers, and indicate actively proliferating cells. The control (R450-treated) graft in the bottom panel of Figure 21 also exhibits evidence of proliferation, including parakeratosis and Ki67-positive cells appearing as brown-staining nuclei. The center panel of Figure 21 exhibits the oligonucleotide #27-treated graft. This graft exhibits significantly reduced proliferation as evidenced by normal (thin) epidermis, lack of invaginations, and substantial loss of Ki67-positive cells.

These results indicate that treatment of human psoriatic grafts with an oligonucleotide targeted to mRNA encoding the IGF-I receptor results in inhibition of epidermal proliferation.

EXAMPLE 21

Topical formulations of complexes of oligonucleotides with cytofectin GSV in aqueous or methylcellulose gel formulations were prepared and assessed foruptake of the oligonucleotide by keratinocytes *in vivo*. The topical formulations contained oligonucleotides complexed with cytofectin GSV in an aqueous solution or methylcellulose carrier, as taught herein. With both aqueous and methylcellulose gel formulations, locatlization of oligonucleotide R451 to nuclei and cytoplasm of keratinocytes in normal human skin grafts on nuce mice was observed. Figure 22 shows an image from confocal microscopy demonstrating oligonucleotide locatlization in the nuclei and cytoplasm of keratinocytes in normal human skin grafts after topical application of fluroescently labeled oligonucleotide (10μM R451) complexed with cytofectin GSV (10μg/ml). Figure 23 shows an image from confocal microscopy demonstrating that topical application of the

same oligonucleotide/GSV concentrations in a 3% (w/v) methylcellulose gel produced similar uptake in the target keratinocyte population. Using an aqueous formulation of oligonucleotide/GSV complexes, penetration of oligonucleotide into the viable epidermis was observed, whereas application of formulations of oliognucleotide complexed with other cationic lipids resulted in localization of oligonucleotide in the stratum corneum.

EXAMPLE 22

Thirteen antisense oligonucleotides targeted to IGFBP-3 were synthesized. The antisense oligonucleotides are C5-propynyl-dU, Dc15 mer phosphorothioate oligodeoxyribonucleotides. Figure 24 attached hereto is a schematic diagram indicating the position of the thirteen oligonucleotides relative to the IGFBP-3 mRNA structure.

These oligonucleotides were screened for their ability to inhibit IGFBP-3 mRNA levels of HaCaT cells in accordance with the teachings herein. HaCaT cells were grown to 90% confluence in DMEM supplemented with 10% (v/v) FCS, then placed in complete keratinocyte serum free medium (KSFM, Gibco), which has a defined amount of EGF, for 24 hours. Oligonucleotides (30nM or 100nM) were complexed with GSV cytofectin (2μg/ml) and added to cells in complete KSFM to allow oligonucleotides to enter the nucleus before removal of EGF. Repeat additions were performed at three hours (in serum free DMEM, which releases the EGF inhibition of IGFBP-3 mRNA) and again after another 24 hours. HaCaT cells were also treated with randomized sequence oligonucleotides (R121, R451, R766 and R961), liposome alone (GSV) or were left untreated (UT). Total RNA was isolated as described in Example 17, 24 hours after the last treatment. Total RNA (15μg) was analyzed by Northern analysis and phosphoroimager quantitation for IGFBP-3 and GADPH mRNA. IGFBP-3 mRNA is expressed as the amount of IGFBP-3 mRNA relative to the amount of GAPDH mRNA.

Figures 25(a)-(d) provide graphs which depict results in this screening process. In these graphs, R1 and R12 refer to R121; R4, R4(0) and R45 rfer to R451; R7, R7(0) and R76 refer to R766; and R9 and R96 refer to R961. The values were standardized to GSV-

treated cells, and data was pooled and statistically analyzed by ANOVA followed by Domet's test to compare each treatment to GSV-treated cells. The pooled data are presented as a bar graph in Figure 26. As demonstrated, at a concentration of 30nM, treatment of HaCaT cells with 8 of the 12 targeted oligonucleotides tested resulted in a statistically significant reduction in levels of IGFBP-3 mRNA relative to GSV-treated cells. At a concentration of 100nM, treatment with 9 fo the 13 targeted oligonucleotides tested resulted in a statistically significant reduction in levels of IGFBP-3 mRNA relative to GSV-treated cells.

These experiments demonstrate that antisense oligonucleotides targeted to genetic sequences encoding IGFBP-3 can inhibit IGFBP-3 mRNA levels in human keratinocytes *in vitro*.

EXAMPLE 23

IGF-I receptor is a potent mitotic signalling molecule for keratinocytes and the human receptor elicits separate intracellular signals that prevent apoptosis (19). It is proposed in accordance with the present invention that inactivation of IGF-I receptors in epidermal keratinocytes will achieve three important outcomes in subsequent UV treatment of lesions:

(i) Acute epidermal hyperplasia following UV has been suggested to increase the risk of keratinocyte carcinogenic transformation (22). By reducing IGF-I receptor expression in the epidermis, the incidence of epidermal hyperplasia following UV exposure is likely to be reduced leading to an overall acceleration in normalization of the lesion and reduced carcinogenic risk.

(ii) Inhibition of anti-apoptotic action of IGF-I receptor will enhance the reversal of epidermal thickening and accelerate normalization of differentiation. Topical or injected IGF-I receptor antisense as adjunctive treatment will increase apoptosis in

the epidermal layer thereby enhancing the reduction in acanthosis observed in UV treatments.

(iii) Survival of keratinocytes, ie. those which evade apoptosis is likely to occur when cells have damaged DNA. Such mutations may be in the tumor suppressor region. Consequently, the use of antisense therapy will result in less frequent selection of mutated keratinocytes and therefore reduced incidence of basal cell carcinomas and squamous.

Accordingly, antisense therapy, especially against IGF-I-receptor is useful in combination with UV therapy in the treatment of epidermal hyperplasia.

EXAMPLE 24

HaCaT cells were treated with antisense oligonucleotides directed to IGF-I receptor mRNA. Levels of IGF-I receptor mRNA were then monitored. In essence, confluent HaCaT cells were treated every 24 hours for four days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM IGF-I receptor specific oligonucleotides (#26 to #86) or random sequence oligonucleotides (*R121, R451 and R766*). Figure 27(a) is a photographic representation showing representative RNase protection assay gel showing IGF-I receptor (IGFR) and GAPDH mRNA in untreated or treated HaCaT cells. Figure 27(b) is a densitometric quantification of IGF-I receptor mRNA in a HaCaT cells following treatment with IGF-I receptor specific oligonucleotides (solid black) random sequence oligonucleotides (horizontal striped bar) or GSV alone (shaded bar) compared to untreated cells (UT, vertical striped bar).

EXAMPLE 25

In this example, reduction in total cellular IGF-I receptor protein was monitored following antisense oligonucleotide treatment. Confluence HaCaT cells were treated with 24 hours for 4 days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM IGF-I receptor specific AONS (#27, #50 and #64) or the random sequence oligonucleotide, R451. Total

cellular protein was isolated and analysed for IGF-I receptor by SDS PAGE followed by western blotting with antibody specific for the human IGF-I receptor. Figure 28(a) shows duplicate treated cellular extracts following the IGF-I receptor at the predicted size of 110 kD. Figure 28(b) is a densitometric quantification of IGF-I receptor protein.

EXAMPLE 26

The reduction in IGF-I receptor numbers was determined on the keratinocyte cell surface after antisense oligonucleotide treatment. HaCaT cells were tranfected with IGF-I receptor specific AONs #27, #50, #64, a random sequence oligonucleotides (R451) or following treatment with GSV a lipid alone every 24 hours for 4 days. Competition binding assays using ^{125}I-IGF-I and the receptor-specific analogue, des(1-3)IGF-I were performed. Results are shown in Figure 29.

EXAMPLE 27

In this example, the apoptotic protecting effects of IGF-I receptor on keratinocyte cells was tested by following the reduction in keratino cell numbers following antisense oligonucleotide treatment. HaCaT cells, initially at 40% confluence, were transfected with the IGF-I receptor specific AON #64, control sequences R451 and 6414 or treated with GSV a lipid alone every 24 hours for 2 days. The cell number was measured in culture wells using a dye binding assay. The results are presented in Figure 30. The results clearly confirm that the IGF-I receptor exhibits an anti-apoptotic effect. By reducing IGF-I receptor levels using antisense oligonucleotide treatment, the anti-apoptotic effect is interrupted and apoptosis results in the reduction in keratinocyte cell number. Results are shown in Figure 30.

EXAMPLE 28

This example shows a reversal of epidermal hyperplasia in psoriatic human skin grafts on nude mice following intradermal injection with antisense oligonucleotides. Grafted psoriasis lesions were injected with IGF-I receptor specific AONs, a random sequence oligonucleotide in PBS, or with PBS alone, every 2 days for 20 days, then analysed

histologically. The results are shown in Figure 31. In Figure 31(a), donor A graft treated with AON #50 showing epidermal thinning compared with the pregraft and control (PBS) treated graft and donor graft treated with AON #27 showing epidermal thinning compared with pregraft and control (R451) treated graft. In Figure 31(b), the mean epidermal cross-sectional area over the full width of grafts is shown as determined by digital image analysis. The results show that epidermal hyperplasia is reversed following the intradermal injection of antisense oligonucleotides.

EXAMPLE 29

Figure 32 shows the reversal of epidermal hyperplasia correlating with reduced IGF-I receptor mRNA in grafted psoriasis lesions treated with antisense oligonucleotides. Figure 32(a) shows a psoriasis lesion prior to grafting and after grafting and treatment with IGF-I receptor specific oligonucleotide #27 (AON #27) or random sequence (R451) immunostained with antibodies to Ki67 to identify proliferating cells. Proliferating cells are indicated by a dark brown nucleus (arrows). Figure 32(b) shows the same lesion prior to grafting and after oligonucleotide treatment as in Figure 32(a) but subjected to *in situ* hybridisation with ^{35}S-labelled cRNA probe complementary to the human IGF-I receptor mRNA. The presence of IGF-I receptor mRNA is indicated by silver grains which are almost eliminated in the epidermis of the lesion treated with IGF-I receptor specific oligonucleotide # 27 (AON #27). This experiment shows that reversal of epidermal hyperplasia correlates with reduced IGF-I receptor mRNA in grafted psoriasis lesions treated with antisense oligonucleotides.

EXAMPLE 30

Figure 33 treatment with oligonucleotides. HaCaT cell monolayers were grown to 90% confluence in DMEM containing 10% fetal calf serum treated every 24 hours for two days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM oligonucleotide. Total RNA was isolated and analysed for IGF-I receptor and GAPDH mRNA using a commercially available ribonuclease protection assay kit. The results show a reduction in IGF-I receptor mRNA in the HaCaT keratinocyte cells.

EXAMPLE 31

Figure 34 treatment with oligonucleotides. HaCaT cell monolayers were grown to 90% confluence in DMEM containing 10% fetal calf serum treated every 24 hours for 4 days with 2 μg/ml GSV lipid alone (GSV) or complexed with 30 nM oligonucleotide. Cells were lysed in a buffer containing 50 mM HEPES, 150 mM NaCl, 10% v/v glycerol, 1 v/v Trison X-100 and 100 μg/ml aprotinin on ice for 30 minutes, then 30 μg of lysate was loaded onto a denaturing 7% w/v polyacrylamide gel followed by transfer onto an Immobilon-P membrane. Membranes were then incubated with anti-IGF-I receptor antibodies C20 (available from Santa Cruz Biotechnology Inc., Santa Cruz, California) for 1 hour at room temperature and developed using the Vistra ECF western blotting kit (Amersham). The results shown in Figure 34 confirm that IGF-I receptor protein is reduced in HaCaT keratinocytes following treatment with oligonucleotides.

EXAMPLE 32

This example shows a reduction in HaCaT keratinocyte cell number following treatment with oligonucleotides. The results are shown in Figure 35. HaCaT cell monolayers were grown at 40% confluence in DMEM containing 10% fetal calf serum treated every 24 hours for 3 days with 2 μg/ml GSV lipid alone (GSV) or complexed with 15 nM oligonucleotide. Cell numbers were then measured every 24 hours using the amido black dye binding assay [32]. Results show that HaCaT keratino cells decrease in number following treatment with oligonucleotides due to a reduction in the anti-apoptotic effect of the IGF-I receptor.

Those skilled in the art will appreciate that the invention described herein is susceptible to variations and modifications other than those specifically described. It is to be understood that the invention includes all such variations and modifications. The invention also includes all of the steps, features, compositions and compounds referred to or indicated in this specification, individually or collectively, and any and all combinations of any two or more of said steps or features.

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CLAIMS:

1. A method for ameliorating the effects of a medical disorder such as a proliferative and/or inflammatory skin disorder in a mammal, said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation or a cell otherwise involved with said medical disorder with an effective amount of a nucleic acid molecule or chemical analogue thereof capable of inhibiting or otherwise reducing growth factor mediated cell proliferation and/or inflammation and/or other medical disorders.

2. A method according to claim 1 wherein cell proliferation and/or inflammation or other medical disorder is mediated by at least one of insulin-like growth factor I (IGF-I), keratinocyte growth factor (KGF), transforming growth factor-α (TGFα), tumour necrosis factor-α (TNFα), interleukin (IL) -1 (IL-1), IL-4, IL-6, IL-8 and/or basic fibroblast growth factor (bFGF).

3. A method according to claim 2 wherein cell proliferation and/or inflammation or other medical disorder is mediated by IGF-I.

4. A method according to claim 1 wherein the nucleic acid molecule inhibits or otherwise reduces IGF-I mediated cell proliferation and/or inflammation or other medical disorder.

5. A method according to claim 1 wherein the proliferative or inflammatory skin disorder is psoriasis, ichthyosis, pityriasis, rubra, pilaris, serborrhoea, keloids, keratosis, neoplasias, scleroderma, warts, benign growths or cancers of the skin.

6. A method according to claim 5 wherein the skin condition is psoriasis.

7. A method according to claim 1 wherein the other medical disorder is a hyperneovascular condition such as a neovascular condition of the retina, brain or skin, growth factor-mediated malignancies, other sclerotic disease, kidney disease or hyperproliferation of the inside of blood vessels or any other hyperplasia.

8. A method according to claim 1 or 4 or 6 or 7 wherein the mammal is a human.

9. A method according to claim 1 or 4 or 6 or 7 wherein the nucleic acid molecule is capable of inhibiting, reducing or otherwise interfering with IGF-I-interaction with its receptor.

10. A method according to claim 9 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGF-I, IGF-I-receptor or an IGF binding protein (IGFBP).

11. A method according to claim 10 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGFBP-2, -3, -4, -5 or -6.

12. A method according to claim 11 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGFBP-2 or IGFBP-3.

13. A method according to claim 10 or 12 wherein the antisense molecule is at least about 15 nucleotides in length and is capable of interacting with at least one sequence selected from the list set forth in Example 6 or Example 7 or Example 8.

14. A method according to claim 12 wherein the antisense molecule comprises the nucleotide sequence:

5'-ATCTCTCCGCTTCCTTTC-3' (<400>10)

15. A method according to claim 12 wherein the antisense molecule is selected from the following:

UCCGGAGCCAGACUU (<400>12)
CACAGUUGCUGCAAG (<400>13)
UCUCCGCUUCCUUUC (<400>14)
AGCCCCCACAGCGAG (<400>15)
GCCUUGGAGAUGAGC (<400>16)
UAACAGAGGUCAGCA (<400>17)
GGAUCAGGGACCAGU (<400>18)
CGGCAAGCUACACAG (<400>19)
GGCAGGCAGGCACAC (<400>20)

16. A method according to claim 15 wherein the antisense molecule in <400>12, <400>13 or <400>14.

17. A method according to claim 15 wherein the antisense molecule in <400>12.

18. A nucleic acid molecule comprising at least about 10 nucleotides capable of hybridising to or forming a heteroduplex or otherwise interacting with a complementary form of <400>12 to <400>20 inclusive.

19. A nucleic acid molecule comprising at least about 15 nucleotides capable of hybridising to or form a heteroduplex or otherwise interacting with a complementary form of <400>12 to <400>14 inclusive.

20. A method of ameliorating the effects of psoriasis or other medical disorder, said method comprising contacting proliferating skin or skin capable of proliferation or cell otherwise associated with said medical disorder with an effective amount of one or more nucleic acid molecules or chemical analogues thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation other medical disorder

wherein said one or more molecules comprises a polynucleotide capable of interacting with mRNA directed from an IGF-I gene, an IGF-I receptor gene or a gene encoding an IGFBP.

21. A method according to claim 20 wherein the IGFBP is IGFBP-2 or IGFBP-3.

22. A method according to claim 20 or 21 wherein the mammal is a human.

23. A method according to claim 22 wherein the nucleic acid molecule is capable of interacting with a nucleotide sequence selected from the list set forth in <400>12 to <400>14 inclusive.

24. A method according to claim 23 wherein the nucleic acid molecule comprises the nucleotide sequence selected from <400>12 to <400>14.

25. A composition comprising a nucleic acid molecule capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation or other medical disorder said composition further comprising one or more pharmaceutically acceptable carriers and/or diluents.

26. A composition according to claim 25 wherein the nucleic acid molecule is antisense molecule to a gene encoding IGF-I, IGF-I-receptor or an IGFBP.

27. A composition according to claim 26 wherein the nucleic acid molecule is selected from <400>12 to <400>20 inclusive.

28. A composition according to claim 26 selected from <400>12 to <400>14 inclusive.

29. A method for ameliorating the effects of a proliferative and/or inflammatory skin disorder such as psoriasis said method comprising contacting the proliferating and/or inflamed skin or skin capable of proliferation and/or inflammation with effective amounts of UV treatment and a nucleic acid molecule or chemical

analogue thereof capable of inhibiting or otherwise reducing IGF-I mediated cell proliferation and/or inflammation.

30. A method according to claim 29 wherein the proliferative or inflammatory skin disorder is psoriasis, ichthyosis, pityriasis, rubra, pilaris, serborrhoea, keloids, keratosis, neoplasias, scleroderma, warts, benign growths or cancers of the skin.

31. A method according to claim 30 wherein the proliferative or inflammatory skin disorder is psoriasis.

32. A method according to claim 29 or 30 or 31 wherein the nucleic acid molecule is capable of inhibiting, reducing or otherwise interfering with IGF-I-interaction with its receptor.

33. A method according to claim 32 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGF-I, IGF-I-receptor or an IGF binding protein (IGFBP).

34. A method according to claim 33 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGFBP-2, -3, -4, -5 or -6.

35. A method according to claim 34 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGFBP-2 or IGFBP-3.

36. A method according to claim 33 wherein the nucleic acid molecule is an antisense molecule capable of reducing expression of a gene encoding IGF-I receptor.

37. A method according to any one of claims 29 to 36 wherein the antisense molecule is at least about 15 nucleotides in length and is capable of interacting with at least one sequence selected from the list set forth in Example 6 or Example 7 or Example 8.

38. A method according to claim 37 wherein the antisense molecule comprises the nucleotide sequence:

5'-ATCTCTCCGCTTCCTTTC-3' (<400>10)

39. A method according to claim 37 wherein the antisense molecule is selected from the following:

UCCGGAGCCAGACUU (<400>12)
CACAGUUGCUGCAAG (<400>13)
UCUCCGCUUCCUUUC (<400>14)
AGCCCCCACAGCGAG (<400>15)
GCCUUGGAGAUGAGC (<400>16)
UAACAGAGGUCAGCA (<400>17)
GGAUCAGGGACCAGU (<400>18)
CGGCAAGCUACACAG (<400>19)
GGCAGGCAGGCACAC (<400>20)

40. A method according to claim 39 wherein the antisense molecule in <400>12, <400>13 or <400>14.

41. A method according to claim 40 wherein the antisense molecule in <400>12.

42. A method according to claim 39 wherein the UV treatment occurs simultaneously with or following contact with the nucleic acid molecule or its chemical analogue.

43. Use of an antisense molecule directed to the gene encoding IGF-I receptor or its mRNA as adjunct therapy in combination with UV treatment to reduce proliferation and/or inflammation of keratinocyte cells.

44. Use according to claim 43 in the treatment of psoriasis.

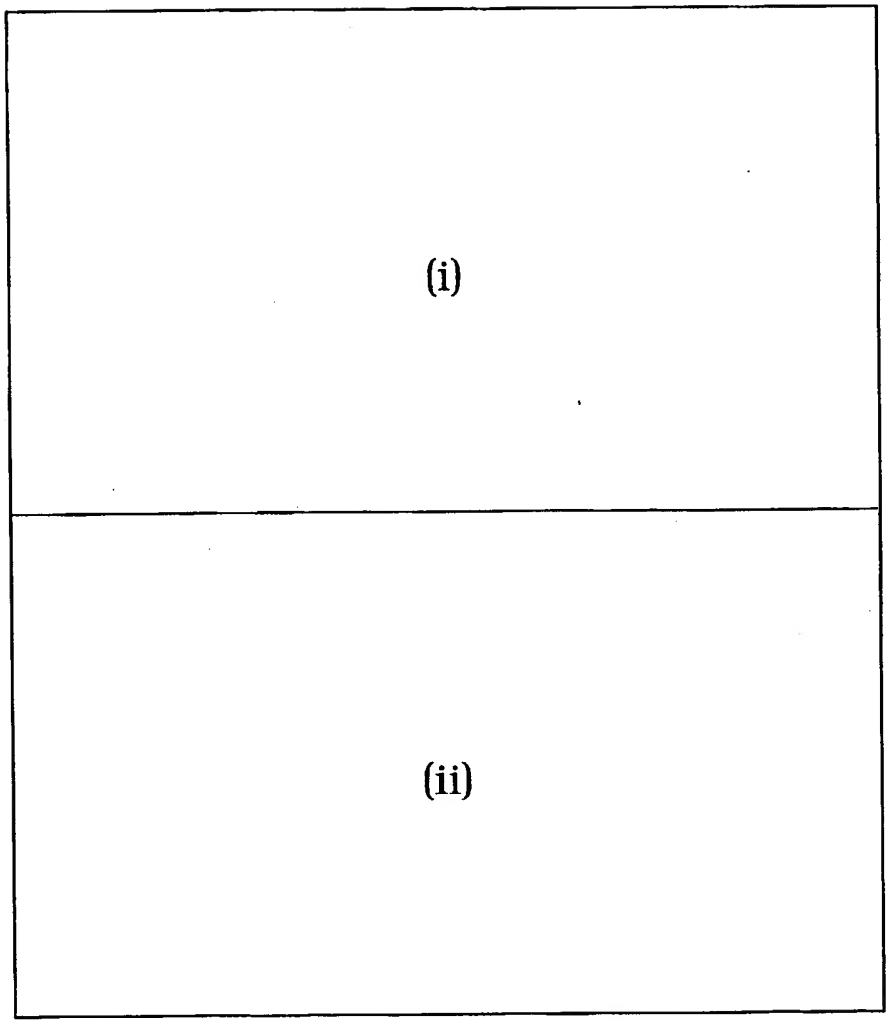


Figure 1

```
1    ATTCGGGGCG AGGGAGGAGG AAGAAGCGGA GGAGGCGGCT CCCGCTCGCA
51   GGGCCGTGCA CCTGCCCGCC CGCCCGCTCG CTCGCTCGCC CGCCGCGCCG
101  CGCTGCCGAC CGCCAGCATG CTGCCGAGAG TGGGCTGCCC CGCGCTGCCG
151  CTGCCGCCGC CGCCGCTGCT GCCGCTGCTG CCGCTGCTGC TGCTGCTACT
201  GGGCGCGAGT GGCGGCGGCG GCGGGGCGCG CGCGGAGGTG CTGTTCCGCT
251  GCCCGCCCTG CACACCCGAG CGCCTGGCCG CCTGCGGGCC CCCGCCGGTT
301  GCGCCGCCCG CCGCGGTGGC CGCAGTGGCC GGAGGCGCCC GCATGCCATG
351  CGCGGAGCTC GTCCGGGAGC CGGGCTGCGG CTGCTGCTCG GTGTGCGCCC
401  GGCTGGAGGG CGAGGCGTGC GGCGTCTACA CCCCGCGCTG CGGCCAGGGG
451  CTGCGCTGCT ATCCCCACCC GGGCTCCGAG CTGCCCCCTGC AGGCGCTGGT
501  CATGGGCGAG GGCACTTGTG AGAAGCGCCG GGACGCCGAG TATGGCGCCA
551  GCCCCGGAGCA GGTTGCAGAC AATGGCGATG ACCACTCAGA AGGAGGCCTG
601  GTGGAGAACC ACGTGGACAG CACCATGAAC ATGTTGGGCG GGGGAGGCAG
651  TGCTGGCCGG AAGCCCCTCA AGTCGGGTAT GAAGGAGCTG GCCGTGTTCC
701  GGGAGAAGGT CACTGAGCAG CACCGGCAGA TGGGCAAGGG TGGGCAAGCAT
```

Figure 1(i)

```
751   CACCTTGGCC TGGAGGAGCC CAAGAAGCTG CGACCACCCC CTGCCAGGAC
801   TCCCTGCCAA CAGGAACTGG ACCAGGTCCT GGAGCGGATC TCCACCATGC
851   GCCTTCCGGA TGAGCGGGGC CCTCTGGAGC ACCTCTACTC CCTGCACATC
901   CCCAACTGTG ACAAGCATGG CCTGTACAAC CTCAAACAGT GCAAGATGTC
951   TCTGAACGGG CAGCGTGGGG AGTGCTGGTG TGTGAACCCC AACACCGGGA
1001  AGCTGATCCA GGGAGCCCCC ACCATCCGGG GGGACCCCGA GTGTCATCTC
1051  TTCTACAATG AGCAGCAGGA GGCTTGCGGG GTGCACACCC AGCGGATGCA
1101  GTAGACCGCA GCCAGCCGGT GCCTGGCGCC CCTGCCCCCC GCCCCTCTCC
1151  AAACACCGGC AGAAAACGGA GAGTGCTTGG GTGGTGGGTG CTGGAGGATT
1201  TTCCAGTTCT GACACACGTA TTTATATTTG GAAAGAGACC AGCACCGAGC
1251  TCGGCACCTC CCCGGCCTCT CTCTTCCCAG CTGCAGATGC CACACCTGCT
1301  CCTTCTTGCT TTCCCCGGGG GAGGAAGGGG GTTGTGGTCG GGGAGCTGGG
1351  GTACAGGTTT GGGGAGGGGG AAGAGAAATT TTTATTTTTG AACCCCTGTG
1401  TCCCTTTTGC ATAAGATTAA AGGAAGGAAA AGT
```

Figure 1(ii)

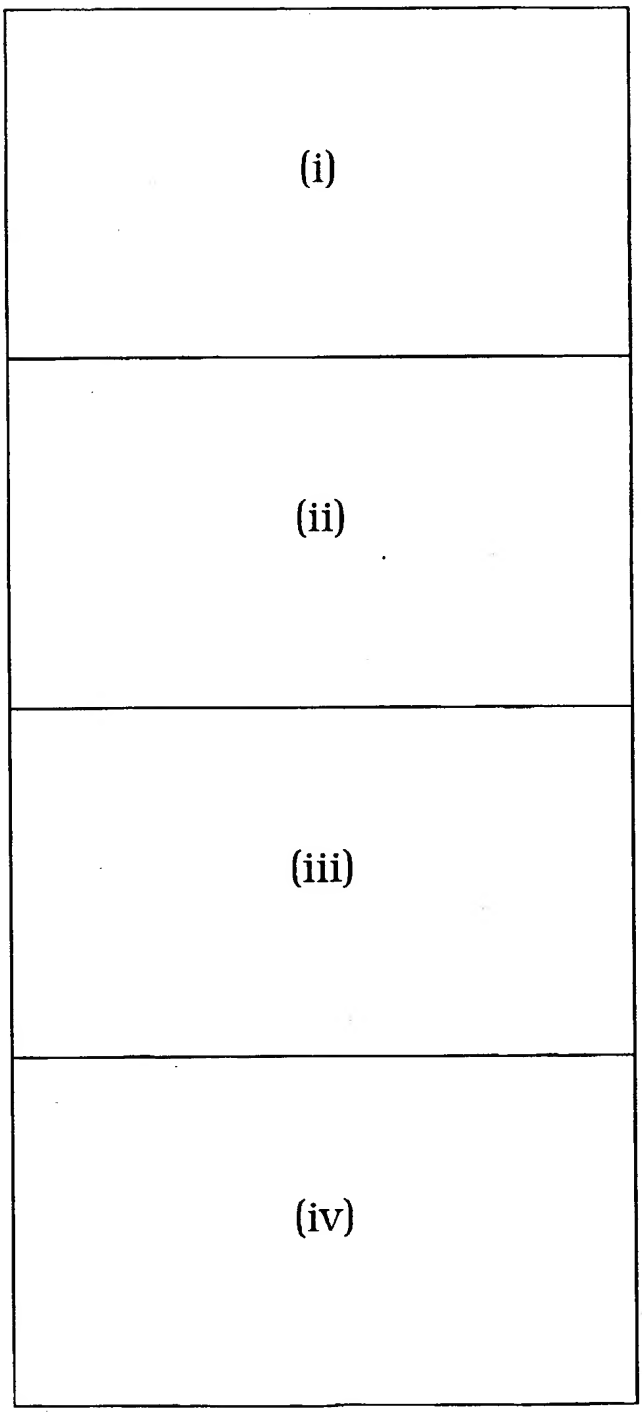


Figure 2

```
1     CTCAGCGCCC AGCCGCTTCC TGCCTGGATT CCACAGCTTC GCGCCGTGTA
51    CTGTCGCCCC ATCCCTGCGC GCCCAGCCTG CCAAGCAGCG TGCCCCGGTT
101   GCAGGCGTCA TGCAGCGGGC GCGACCCACG CTCTGGGCCG CTGCGCTGAC
151   TCTGCTGGTG CTGCTCCGCG GGCCGCCGGT GGCGCGGGCT GGCGCGAGCT
201   CGGGGGGCTT GGGTCCCGTG GTGCGCTGCG AGCCGTGCGA CGCGCGTGCA
251   CTGGCCCAGT GCGCGCCTCC GCCCGCCGTG TGCGCGGAGC TGGTGCGCGA
301   GCCGGGCTGC GGCTGCTGCC TGACGTGCGC ACTGAGCGAG GGCCAGCCGT
351   GCGGCATCTA CACCGAGCGC TGTGGCTCCG GCCTTCGCTG CCAGCCGTCG
401   CCCGACGAGG CGCGACCGCT GCAGGCGCTG CTGGACGGCC GCGGGCTCTG
451   CGTCAACGCT AGTGCCGTCA GCCGCCTGCG CGCCTACCTG CTGCCAGCGC
501   CGCCAGCTCC AGGAAATGCT AGTGAGTCGG AGGAAGACCG CAGCGCCGGC
551   AGTGTGGAGA GCCCGTCCGT CTCCAGCACG CACCGGGTGT CTGATCCCAA
601   GTTCCACCCC CTCCATTCAA AGATAATCAT CATCAAGAAA GGGCATGCTA
651   AAGACAGCCA GCGCTACAAA GTTGACTACG AGTCTCAGAG CACAGATACC
701   CAGAACTTCT CCTCCGAGTC CAAGCGGGAG ACAGAATATG GTCCCTGCCG
```

Figure 2(i)

```
751   TAGAGAAATG GAAGACACAC TGAATCACCT GAAGTTCCTC AATGTGCTGA
801   GTCCCAGGGG TGTACACATT CCCAACTGTG ACAAGAAGGG ATTTTATAAG
851   AAAAAGCAGT GTCGCCCTTC CAAAGGCAGG AAGCGGGGCT TCTGCTGGTG
901   TGTGGATAAG TATGGGCAGC CTCTCCCAGG CTACACCACC AAGGGGAAGG
951   AGGACGTGCA CTGCTACAGC ATGCAGAGCA AGTAGACGCC TGCCGCAAGT
1001  TAATGTGGAG CTCAAATATG CCTTATTTTG CACAAAAGAC TGCCAAGGAC
1051  ATGACCAGCA GCTGGCTACA GCCTCGATTT ATATTTCTGT TTGTGGTGAA
1101  CTGATTTTTT TTAAACCAAA GTTTAGAAAG AGGTTTTTGA AATGCCTATG
1151  GTTTCTTTGA ATGGTAAACT TGAGCATCTT TTCACTTTCC AGTAGTCAGC
1201  AAAGAGCAGT TTGAATTTTC TTGTCGCTTC CTATCAAAAT ATTCAGAGAC
1251  TCGAGCACAG CACCCAGACT TCATGCGCCC GTGGAATGCT CACCACATGT
1301  TGGTCGAAGC GGCCGACCAC TGACTTTGTG ACTTAGGCGG CTGTGTTGCC
1351  TATGTAGAGA ACACGCTTCA CCCCCACTCC CCGTACAGTG CGCACAGGCT
1401  TTATCGAGAA TAGGAAAACC TTTAAACCCC GGTCATCCGG ACATCCCAAC
1451  GCATGCTCCT GGAGCTCACA GCCTTCTGTG GTGTCATTTC TGAAACAAGG
```

Figure 2(ii)

```
1501  GCGTGGATCC CTCAACCAAG AAGAATGTTT ATGTCTTCAA GTGACCTGTA
1551  CTGCTTGGGG ACTATTGGAG AAAATAAGGT GGAGTCCTAC TTGTTTAAAA
1601  AATATGTATC TAAGAATGTT CTAGGGCACT CTGGGAACCT ATAAAGGCAG
1651  GTATTTCGGG CCCTCCTCTT CAGGAATCTT CCTGAAGACA TGGCCCAGTC
1701  GAAGGCCCAG GATGGCTTTT GCTGCGGCCC CGTGGGGTAG GAGGGACAGA
1751  GAGACGGGAG AGTCAGCCTC CACATTCAGA GGCATCACAA GTAATGGCAC
1801  AATTCTTCGG ATGACTGCAG AAAATAGTGT TTTGTAGTTC AACAACTCAA
1851  GACGAAGCTT ATTTCTGAGG ATAAGCTCTT TAAAGGCAAA GCTTTATTTT
1901  CATCTCTCAT CTTTTGTCCT CCTTAGCACA ATGTAAAAAA GAATAGTAAT
1951  ATCAGAACAG GAAGGAGGAA TGGCTTGCTG GGGAGCCCAT CCAGGACACT
2001  GGGAGCACAT AGAGATTCAC CCATGTTTGT TGAACTTAGA GTCATTCTCA
2051  TGCTTTTCTT TATAATTCAC ACATATATGC AGAGAAGATA TGTTCTTGTT
2101  AACATTGTAT ACAACATAGC CCCAAATATA GTAAGATCTA TACTAGATAA
2151  TCCTAGATGA AATGTTAGAG ATGCTATATG ATACAACTGT GGCCATGACT
2201  GAGGAAAGGA GCTCACGCCC AGAGACTGGG CTGCTCTCCC GGAGGCCAAA
```

Figure 2(iii)

```
2251  CCCAAGAAGG TCTGGCAAAG TCAGGCTCAG GGAGACTCTG CCCTGCTGCA
2301  GACCTCGGTG TGGACACACG CTGCATAGAG CTCTCCTTGA AAACAGAGGG
2351  GTCTCAAGAC ATTCTGCCTA CCTATTAGCT TTTCTTTATT TTTTTAACTT
2401  TTTGGGGGGA AAAGTATTTT TGAGAAGTTT GTCTTGCAAT GTATTTATAA
2451  ATAGTAAATA AAGTTTTTAC CATT
```

Figure 2(iv)

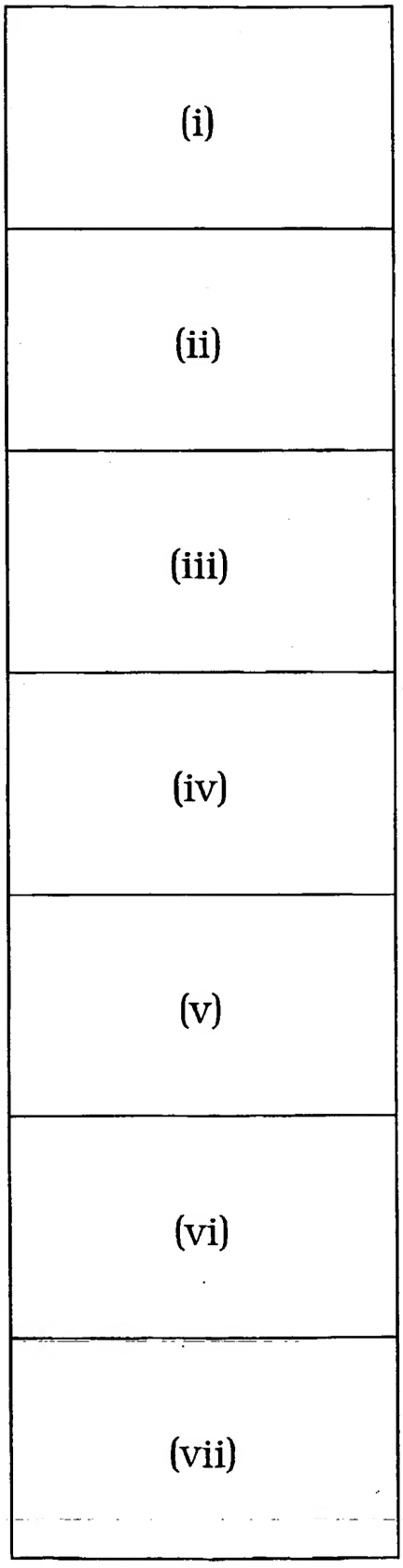


Figure 3

```
1    TTTTTTTTTT TTTTGAGAAA GGGAATTTCA TCCCAAATAA AAGGAATGAA
51   GTCTGGCTCC GGAGGAGGGT CCCCGACCTC GCTGTGGGGG CTCCTGTTTC
101  TCTCCGCCGC GCTCTCGCTC TGGCCGACGA GTGGAGAAAT CTGCGGGCCA
151  GGCATCGACA TCCGCAACGA CTATCAGCAG CTGAAGCGCC TGGAGAACTG
201  CACGGTGATC GAGGGCTACC TCCACATCCT GCTCATCTCC AAGGCCGAGG
251  ACTACCGCAG CTACCGCTTC CCCAAGCTCA CGGTCATTAC CGAGTACTTG
301  CTGCTGTTCC GAGTGGCTGG CCTCGAGAGC CTCGGAGACC TCTTCCCCAA
351  CCTCACGGTC ATCCGCGGCT GGAAACTCTT CTACAACTAC GCCCTGGTCA
401  TCTTCGAGAT GACCAATCTC AAGGATATTG GGCTTTACAA CCTGAGGAAC
451  ATTACTCGGG GGGCCATCAG GATTGAGAAA AATGCTGACC TCTGTTACCT
501  CTCCACTGTG GACTGGTCCC TGATCCTGGA TGCGGTGTCC AATAACTACA
551  TTGTGGGGAA TAAGCCCCCA AAGGAATGTG GGGACCTGTG TCCAGGGACC
601  ATGGAGGAGA AGCCGATGTG TGAGAAGACC ACCATCAACA ATGAGTACAA
651  CTACCGCTGC TGGACCACAA ACCGCTGCCA GAAAATGTGC CCAAGCACGT
701  GTGGGAAGCG GGCGTGCACC GAGAACAATG AGTGCTGCCA CCCCGAGTGC
```

Figure 3(i)

```
751  CTGGGCAGCT GCAGCGCGCC TGACAACGAC ACGGCCTGTG TAGCTTGCCG
801  CCACTACTAC TATGCCGGTG TCTGTGTGCC TGCCTGCCCG CCCAACACCT
851  ACAGGTTTGA GGGCTGGCGC TGTGTGGACC GTGACTTCTG CGCCAACATC
901  CTCAGCGCCG AGAGCAGCGA CTCCGAGGGG TTTGTGATCC ACGACGGCGA
951  GTGCATGCAG GAGTGCCCCT CGGGCTTCAT CCGCAACGGC AGCCAGAGCA
1001 TGTACTGCAT CCCTTGTGAA GGTCCTTGCC CGAAGGTCTG TGAGGAAGAA
1051 AAGAAAACAA AGACCATTGA TTCTGTTACT TCTGCTCAGA TGCTCCAAGG
1101 ATGCACCATC TTCAAGGGCA ATTTGCTCAT TAACATCCGA CGGGGGAATA
1151 ACATTGCTTC AGAGCTGGAG AACTTCATGG GGCTCATCGA GGTGGTGACG
1201 GGCTACGTGA AGATCCGCCA TTCTCATGCC TTGGTCTCCT TGTCCTTCCT
1251 AAAAAACCTT CGCCTCATCC TAGGAGAGGA GCAGCTAGAA GGGAATTACT
1301 CCTTCTACGT CCTCGACAAC CAGAACTTGC AGCAACTGTG GGACTGGGAC
1351 CACCGCAACC TGACCATCAA AGCAGGGAAA ATGTACTTTG CTTTCAATCC
1401 CAAATTATGT GTTTCCGAAA TTTACCGCAT GGAGGAAGTG ACGGGGACTA
1451 AAGGGCGCCA AAGCAAAGGG GACATAAACA CCAGGAACAA CGGGGAGAGA
```

Figure 3(ii)

```
1501  GCCTCCTGTG AAAGTGACGT CCTGCATTTC ACCTCCACCA CCACGTCGAA
1551  GAATCGCATC ATCATAACCT GGCACCGGTA CCGGCCCCCT GACTACAGGG
1601  ATCTCATCAG CTTCACCGTT TACTACAAGG AAGCACCCTT TAAGAATGTC
1651  ACAGAGTATG ATGGGCAGGA TGCCTGCGGC TCCAACAGCT GGAACATGGT
1701  GGACGTGGAC CTCCCGCCCA ACAAGGACGT GGAGCCCGGC ATCTTACTAC
1751  ATGGGCTGAA GCCCTGGACT CAGTACGCCG TTTACGTCAA GGCTGTGACC
1801  CTCACCATGG TGGAGAACGA CCATATCCGT GGGGCCAAGA GTGAGATCTT
1851  GTACATTCGC ACCAATGCTT CAGTTCCTTC CATTCCCTTG GACGTTCTTT
1901  CAGCATCGAA CTCCTCTTCT CAGTTAATCG TGAAGTGGAA CCCTCCCTCT
1951  CTGCCCAACG GCAACCTGAG TTACTACATT GTGCGCTGGC AGCGGCAGCC
2001  TCAGGACGGC TACCTTTACC GGCACAATTA CTGCTCCAAA GACAAAATCC
2051  CCATCAGGAA GTATGCCGAC GGCACCATCG ACATTGAGGA GGTCACAGAG
2101  AACCCCAAGA CTGAGGTGTG TGGTGGGGAG AAAGGGCCTT GCTGCGCCTG
2151  CCCCAAAACT GAAGCCGAGA AGCAGGCCGA GAAGGAGGAG GCTGAATACC
2201  GCAAAGTCTT TGAGAATTTC CTGCACAACT CCATCTTCGT GCCCAGACCT
```

Figure 3(iii)

```
2251  GAAAGGAAGC GGAGAGATGT CATGCAAGTG GCCAACACCA CCATGTCCAG
2301  CCGAAGCAGG AACACCACGG CCGCAGACAC CTACAACATC ACCGACCCGG
2351  AAGAGCTGGA GACAGAGTAC CCTTTCTTTG AGAGCAGAGT GGATAACAAG
2401  GAGAGAACTG TCATTTCTAA CCTTCGGCCT TTCACATTGT ACCGCATCGA
2451  TATCCACAGC TGCAACCACG AGGCTGAGAA GCTGGGCTGC AGCGCCTCCA
2501  ACTTCGTCTT TGCAAGGACT ATGCCCGCAG AAGGAGCAGA TGACATTCCT
2551  GGGCCAGTGA CCTGGGAGCC AAGGCCTGAA AACTCCATCT TTTTAAAGTG
2601  GCCGGAACCT GAGAATCCCA ATGGATTGAT TCTAATGTAT GAAATAAAAT
2651  ACGGATCACA AGTTGAGGAT CAGCGAGAAT GTGTGTCCAG ACAGGAATAC
2701  AGGAAGTATG GAGGGGCCAA GCTAAACCGG CTAAACCCGG GGAACTACAC
2751  AGCCCGGATT CAGGCCACAT CTCTCTCTGG GAATGGGTCG TGGACAGATC
2801  CTGTGTTCTT CTATGTCCAG GCCAAAACAG GATATGAAAA CTTCATCCAT
2851  CTGATCATCG CTCTGCCCGT CGCTGTCCTG TTGATCGTGG GAGGGTTGGT
2901  GATTATGCTG TACGTCTTCC ATAGAAAGAG AAATAACAGC AGGCTGGGGA
2951  ATGGAGTGCT GTATGCCTCT GTGAACCCGG AGTACTTCAG CGCTGCTGAT
```

Figure 3(iv)

```
3001  GTGTACGTTC CTGATGAGTG GGAGGTGGCT CGGGAGAAGA TCACCATGAG
3051  CCGGGAACTT GGGCAGGGGT CGTTTGGGAT GGTCTATGAA GGAGTTGCCA
3101  AGGGTGTGGT GAAAGATGAA CCTGAAACCA GAGTGGCCAT TAAAACAGTG
3151  AACGAGGCCG CAAGCATGCG TGAGAGGATT GAGTTTCTCA ACGAAGCTTC
3201  TGTGATGAAG GAGTTCAATT GTCACCATGT GGTGCGATTG CTGGGTGTGG
3251  TGTCCCAAGG CCAGCCAACA CTGGTCATCA TGGAACTGAT GACACGGGGC
3301  GATCTCAAAA GTTATCTCCG GTCTCTGAGG CCAGAAATGG AGAATAATCC
3351  AGTCCTAGCA CCTCCAAGCC TGAGCAAGAT GATTCAGATG GCCGGAGAGA
3401  TTGCAGACGG CATGGCATAC CTCAACGCCA ATAAGTTCGT CCACAGAGAC
3451  CTTGCTGCCC GGAATTGCAT GGTAGCCGAA GATTTCACAG TCAAAATCGG
3501  AGATTTTGGT ATGACGCGAG ATATCTATGA GACAGACTAT TACCGGAAAG
3551  GAGGCAAAGG GCTGCTGCCC GTGCGCTGGA TGTCTCCTGA GTCCCTCAAG
3601  GATGGAGTCT TCACCACTTA CTCGGACGTC TGGTCCTTCG GGGTCGTCCT
3651  CTGGGAGATC GCCACACTGG CCGAGCAGCC CTACCAGGGC TTGTCCAACG
3701  AGCAAGTCCT TCGCTTCGTC ATGGAGGGCG GCCTTCTGGA CAAGCCAGAC
```

Figure 3(v)

```
3751  AACTGTCCTG ACATGCTGTT TGAACTGATG CGCATGTGCT GGCAGTATAA
3801  CCCCAAGATG AGGCCTTCCT TCCTGGAGAT CATCAGCAGC ATCAAAGAGG
3851  AGATGGAGCC TGGCTTCCGG GAGGTCTCCT TCTACTACAG CGAGGAGAAC
3901  AAGCTGCCCG AGCCGGAGGA GCTGGACCTG GAGCCAGAGA ACATGGAGAG
3951  CGTCCCCCTG GACCCCTCGG CCTCCTCGTC CTCCCTGCCA CTGCCCGACA
4001  GACACTCAGG ACACAAGGCC GAGAACGGCC CCGGCCCTGG GGTGCTGGTC
4051  CTCCCGCGCCA GCTTCGACGA GAGACAGCCT TACGCCCACA TGAACGGGGG
4101  CCGCAAGAAC GAGCGGGCCT TGCCGCTGCC CCAGTCTTCG ACCTGCTGAT
4151  CCTTGGATCC TGAATCTGTG CAAACAGTAA CGTGTGCGCA CGCGCAGCGG
4201  GGTGGGGGGG GAGAGAGAGT TTTAACAATC CATTCACAAG CCTCCTGTAC
4251  CTCAGTGGAT CTTCAGTTCT GCCCTTGCTG CCCGCGGGAG ACAGCTTCTC
4301  TGCAGTAAAA CACATTTGGG ATGTTCCTTT TTTCAATATG CAAGCAGCTT
4351  TTTATTCCCT GCCCAAACCC TTAACTGACA TGGGCCTTTA AGAACCTTAA
4401  TGACAACACT TAATAGCAAC AGAGCACTTG AGAACCAGTC TCCTCACTCT
4451  GTCCCTGTCC TTCCCTGTTC TCCCTTTCTC TCTCCTCTCT GCTTCATAAC
```

Figure 3(vi)

```
4501  GGAAAAATAA TTGCCACAAG TCCAGCTGGG AAGCCCTTTT TATCAGTTTG
4551  AGGAAGTGGC TGTCCCTGTG GCCCCATCCA ACCACTGTAC ACACCCGCCT
4601  GACACCGTGG GTCATTACAA AAAAACACGT GGAGATGGAA ATTTTTACCT
4651  TTATCTTTCA CCTTTCTAGG GACATGAAAT TTACAAAGGG CCATCGTTCA
4701  TCCAAGGCTG TTACCATTTT AACGCTGCCT AATTTTGCCA AAATCCTGAA
4751  CTTTCTCCCT CATCGGCCCG GCGCTGATTC CTCGTGTCCG GAGGCATGGG
4801  TGAGCATGGC AGCTGGTTGC TCCATTTGAG AGACACGCTG GCGACACACT
4851  CCGTCCATCC GACTGCCCCT GCTGTGCTGC TCAAGGCCAC AGGCACACAG
4901  GTCTCATTGC TTCTGACTAG ATTATTATTT GGGGGAACTG GACACAATAG
4951  GTCTTTCTCT CAGTGAAGGT GGGGAGAAGC TGAACCGGC
```

Figure 3 (vii)

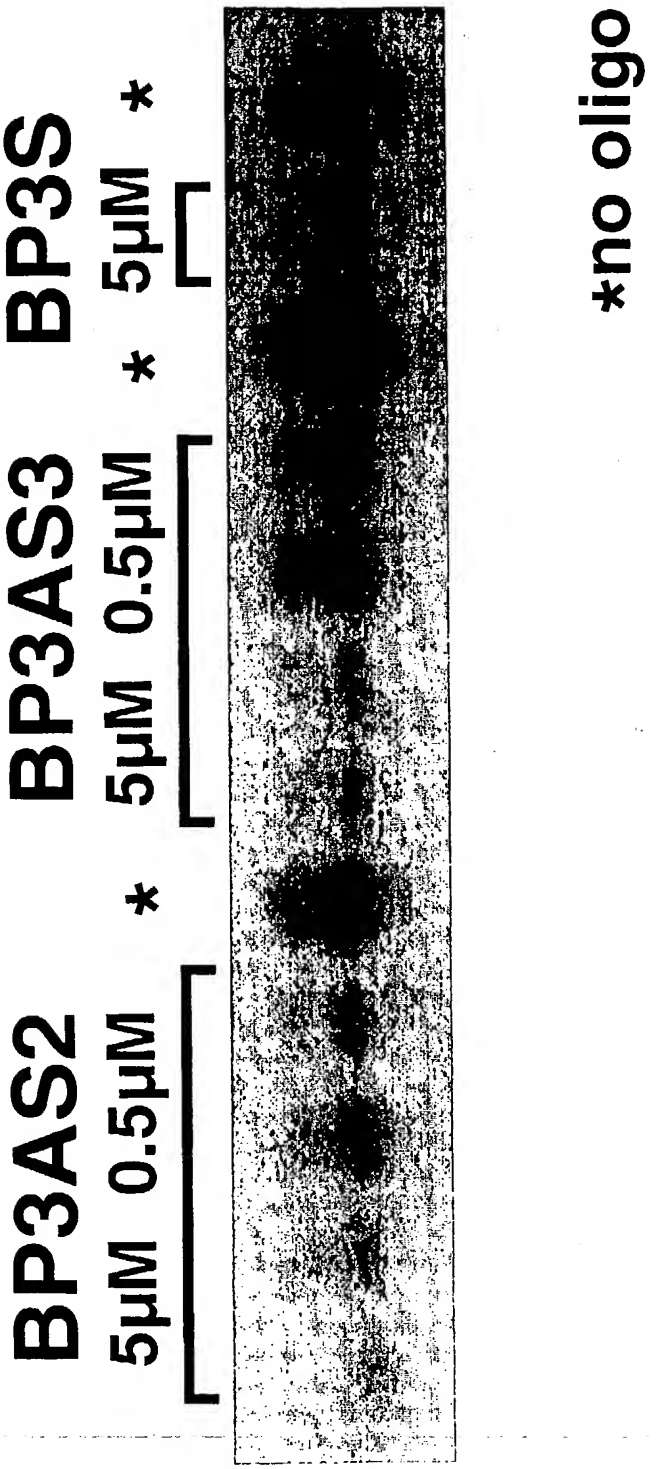


Figure 4a

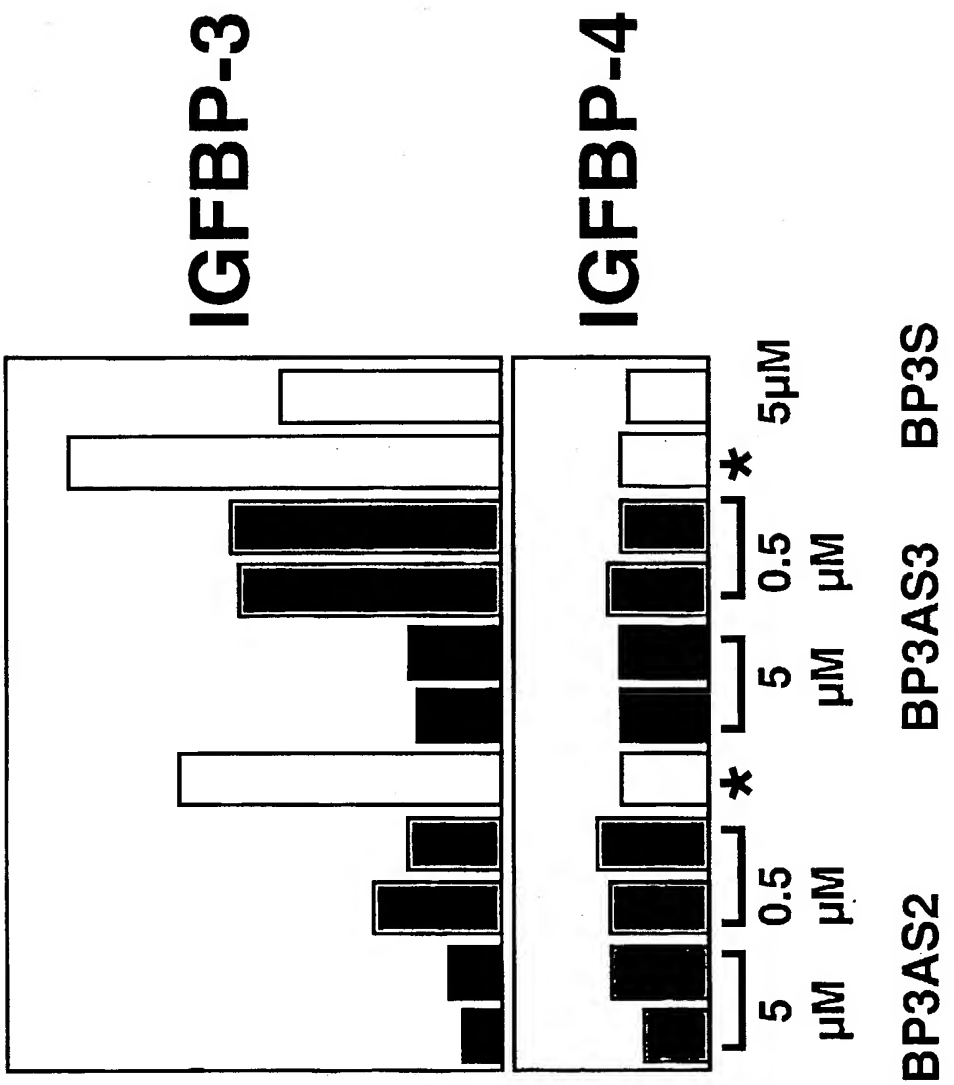


Figure 4b

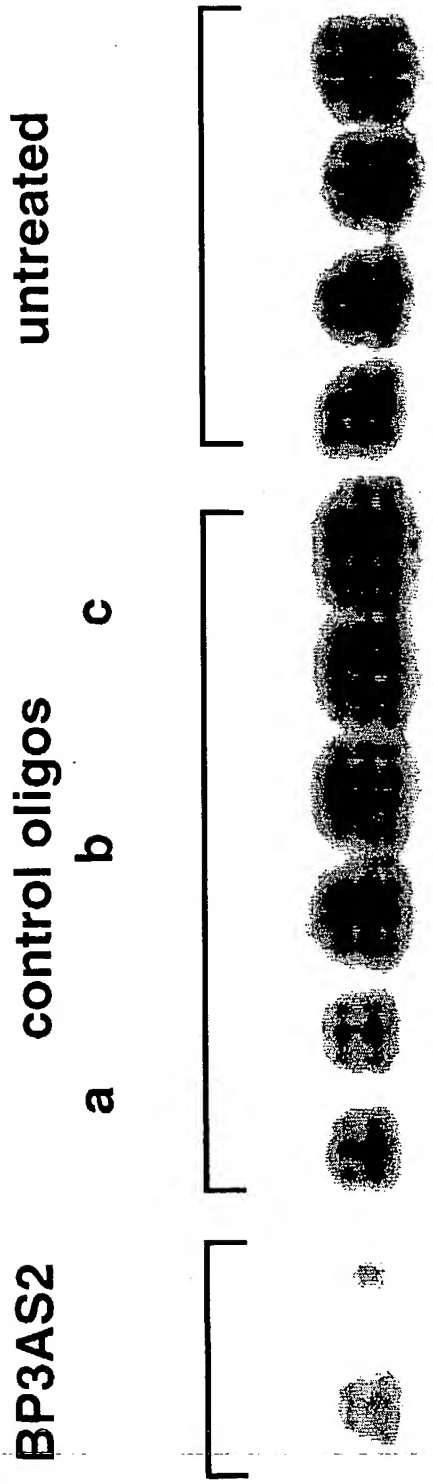


Figure 5a

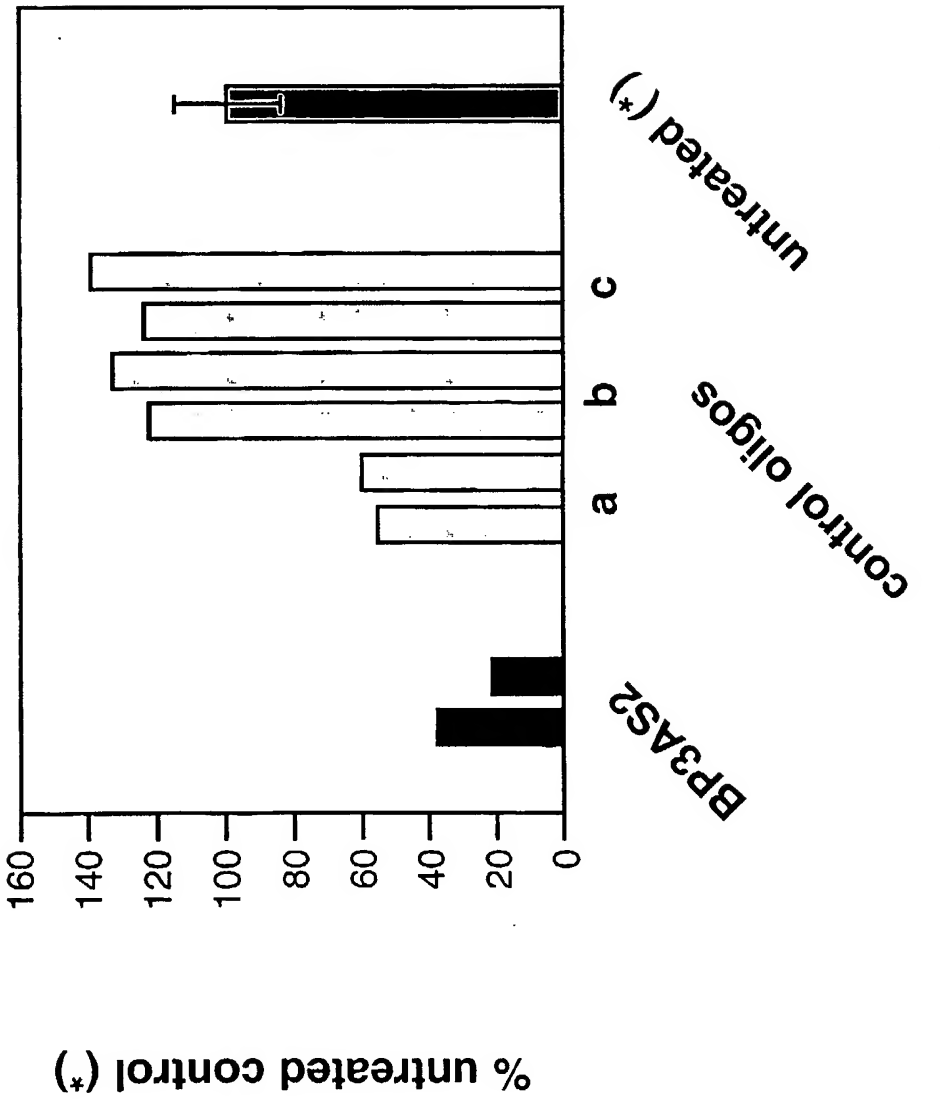

% untreated control (*)
160
140
120
100
80
60
40
20
0
BP3AS2
control oligos
a
b
c
untreated (*)

Figure 5b

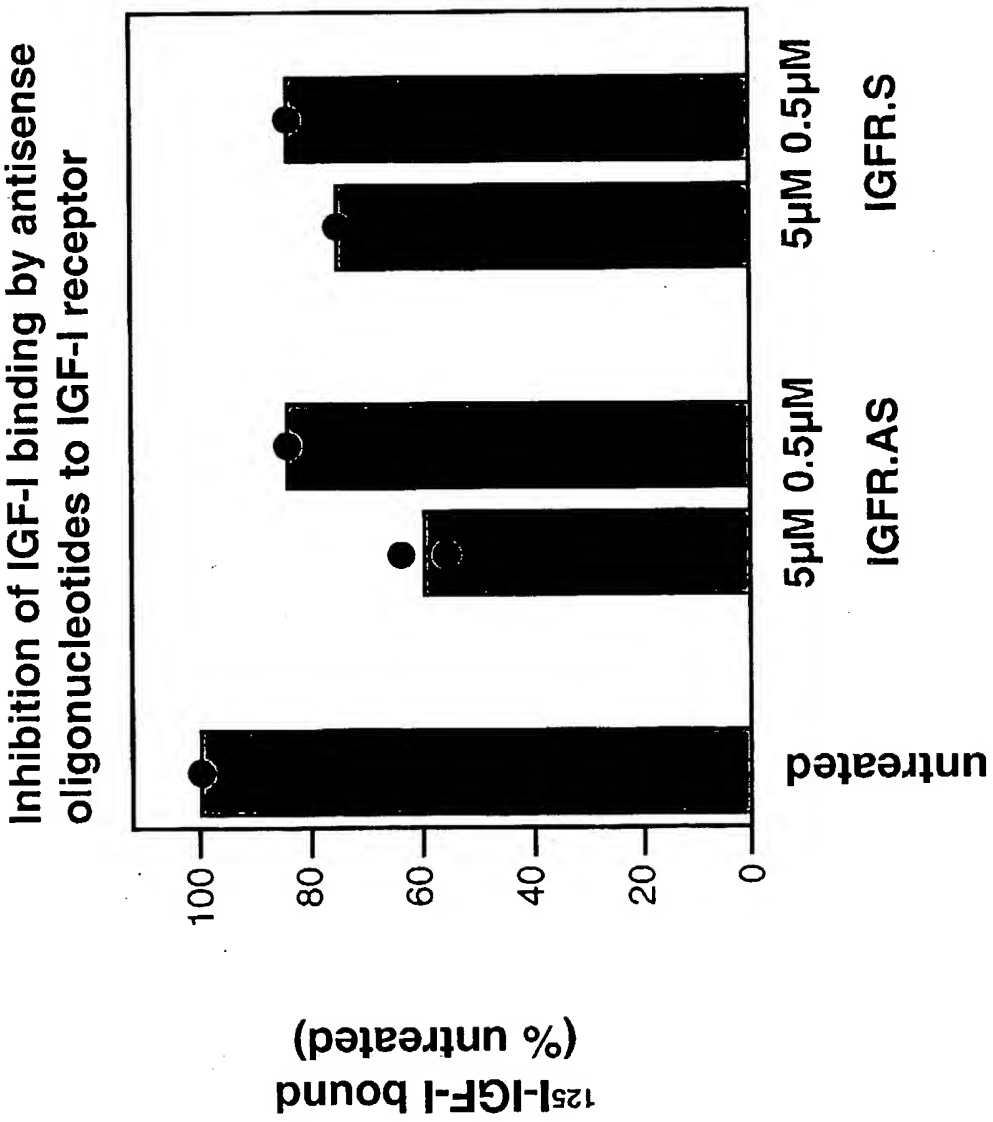

Inhibition of IGF-I binding by antisense oligonucleotides to IGF-I receptor
125I-IGF-I bound (% untreated)
100
80
60
40
20
0
untreated
5μM 0.5μM
IGFR.AS
5μM 0.5μM
IGFR.S

Figure 6

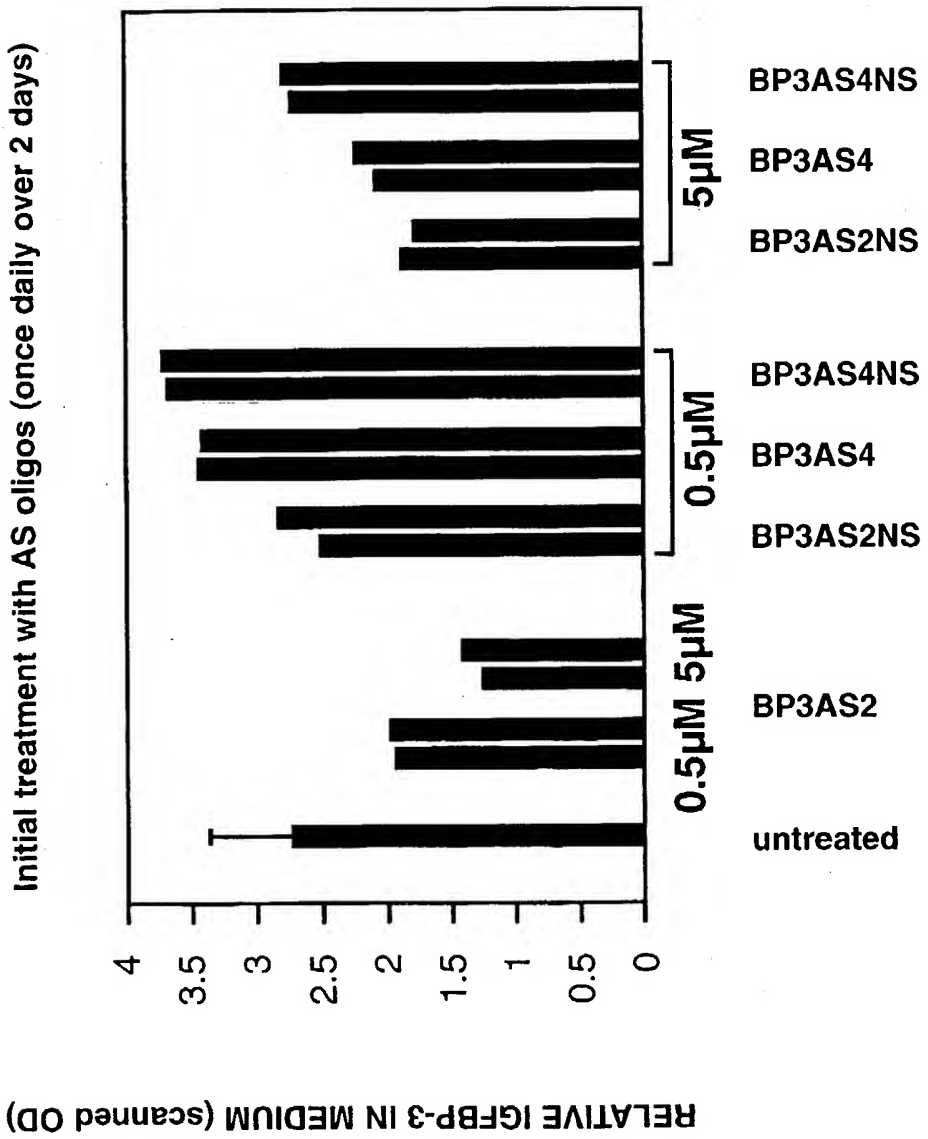

RELATIVE IGFBP-3 IN MEDIUM (scanned OD)
4
3.5
3
2.5
2
1.5
1
0.5
0
Initial treatment with AS oligos (once daily over 2 days)
0.5µM 5µM
untreated
BP3AS2
0.5µM
BP3AS2NS
BP3AS4
BP3AS4NS
5µM
BP3AS2NS
BP3AS4
BP3AS4NS

Figure 7

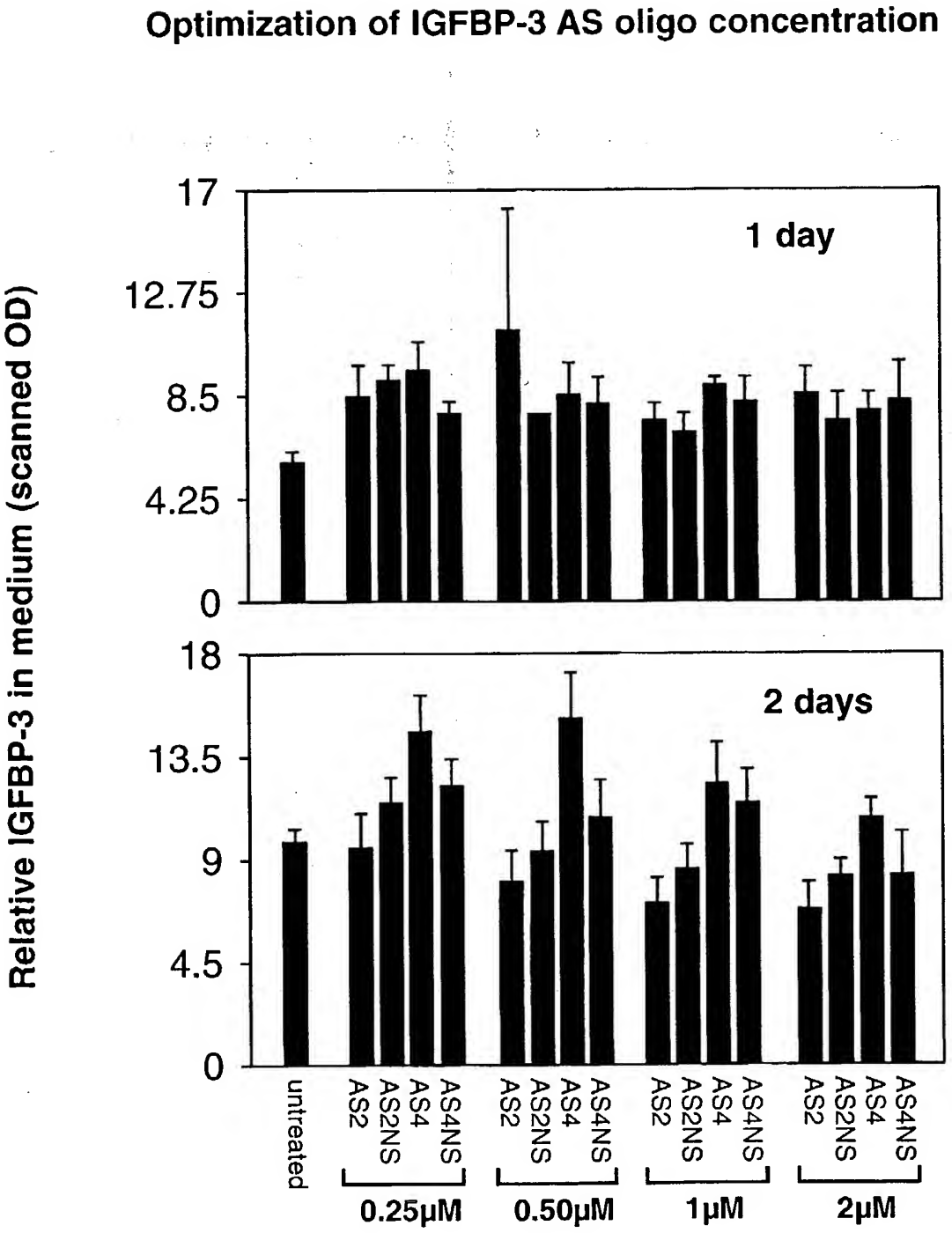

Optimization of IGFBP-3 AS oligo concentration
Relative IGFBP-3 in medium (scanned OD)
17
12.75
8.5
4.25
0
1 day
18
13.5
9
4.5
0
2 days
untreated
AS2
AS2NS
AS4
AS4NS
0.25μM
0.50μM
1μM
2μM

Figure 8

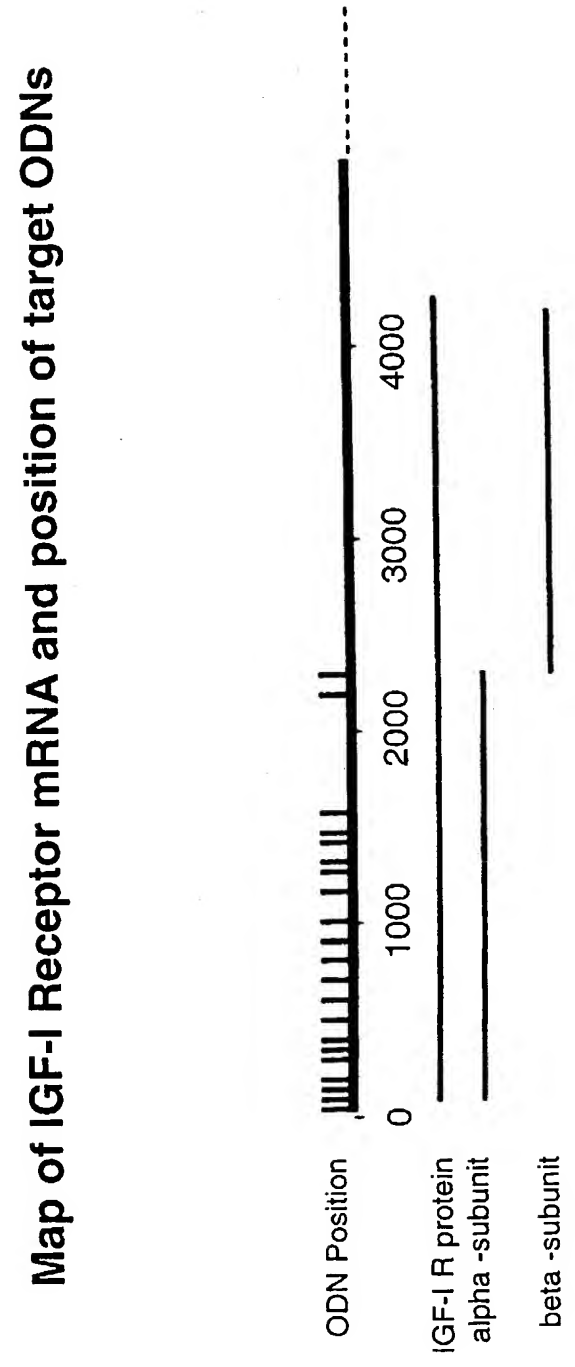
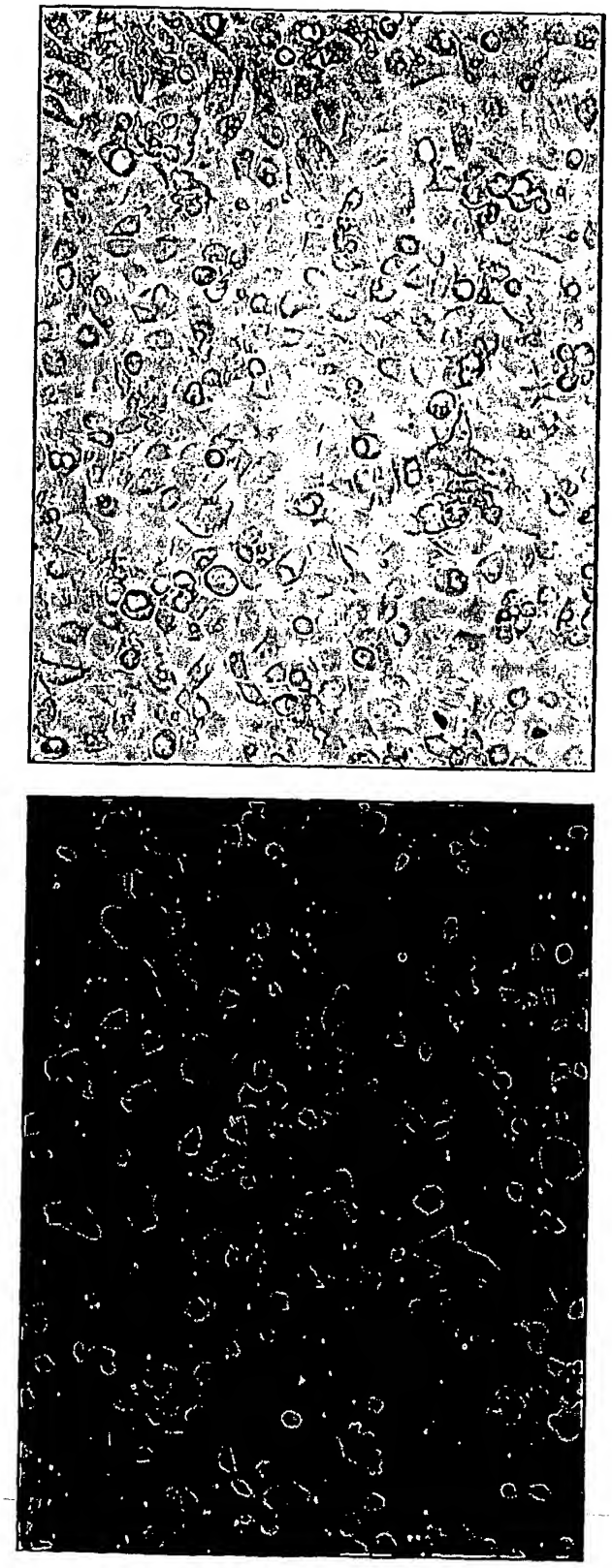

Map of IGF-I Receptor mRNA and position of target ODNs
ODN Position
0
1000
2000
3000
4000
IGF-I R protein
alpha -subunit
beta -subunit
•Position of the 21 tested ODNs (|)
•mRNA transcript lengths = 7Kb and 11Kb
•coding sequence 46-4149

Figure 9

Lipid-mediated uptake of oligonucleotide in keratinocytes

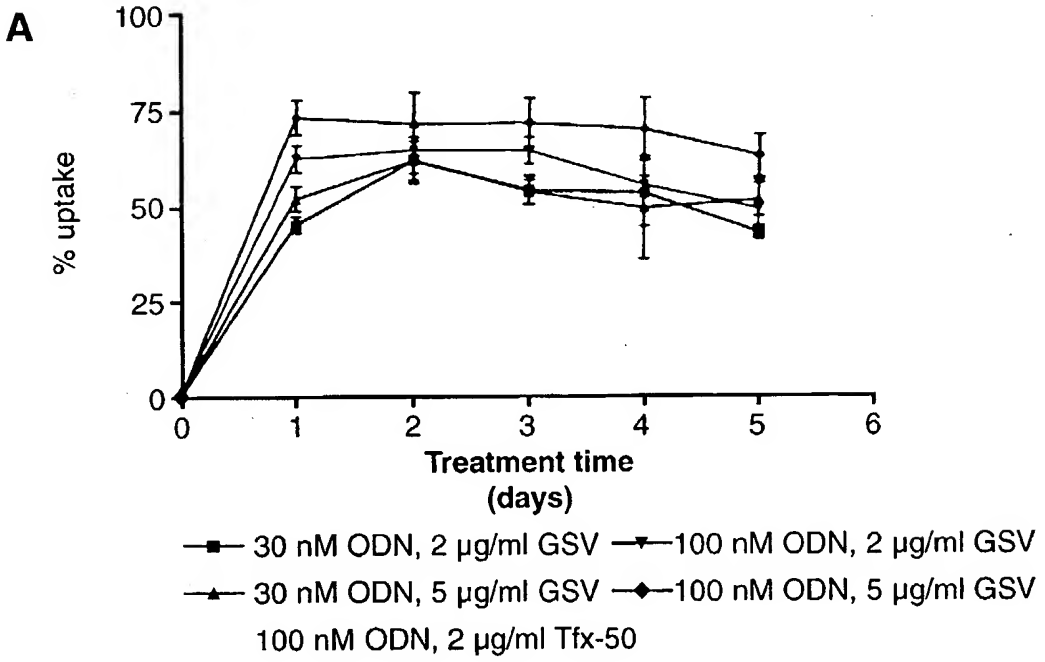


B

A

Figure 10

Uptake (A) and toxicity (B) of ODN/ lipid complexes in keratinocytes



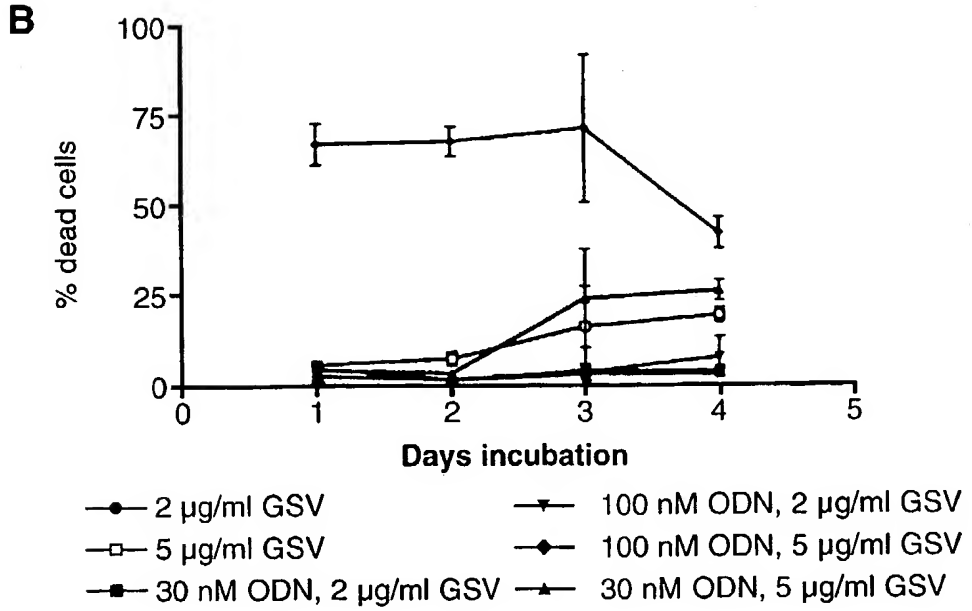
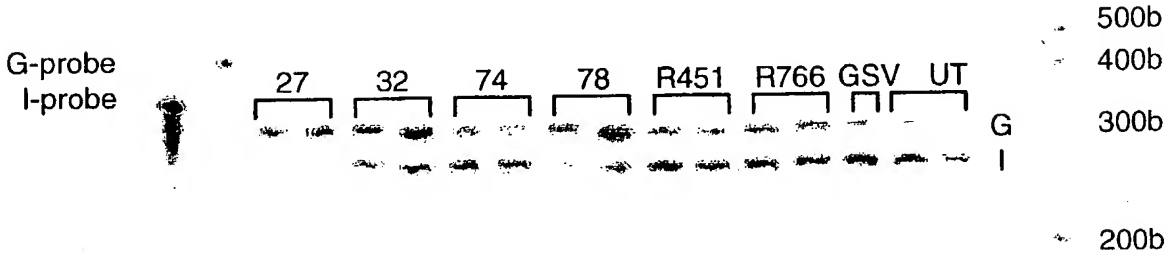


Figure 11

IGF-I Receptor mRNA in ODN treated (30nM) HaCaT cells (2µg/ml GSV)

A



B

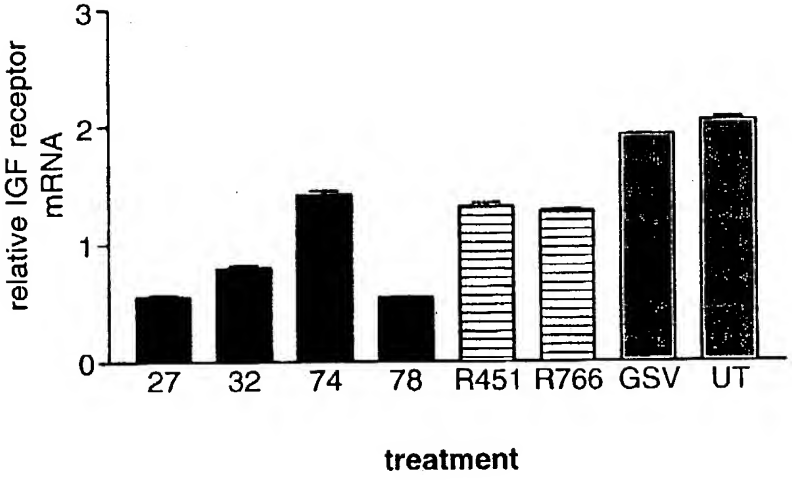


Figure 12

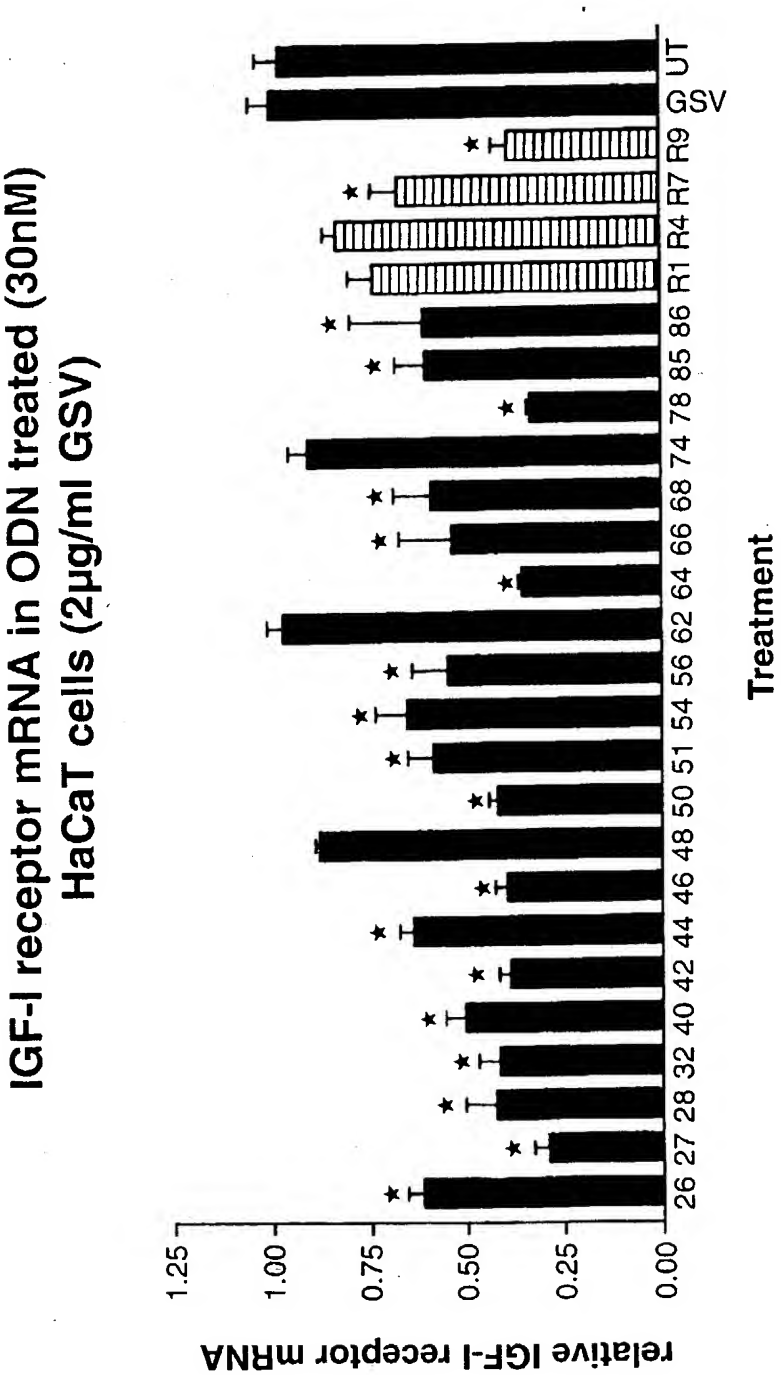

IGF-I receptor mRNA in ODN treated (30nM)
HaCaT cells (2µg/ml GSV)
relative IGF-I receptor mRNA
1.25
1.00
0.75
0.50
0.25
0.00
26 27 28 32 40 42 44 46 48 50 51 54 56 62 64 66 68 74 78 85 86 R1 R4 R7 R9 GSV UT
Treatment

Figure 13

Effect of antisense oligonucleotides on IGF-1 receptor levels on the surface of keratinocytes:
Competition Assay - ^{125}I IGF-1 vs Des 1-3 IGF-1

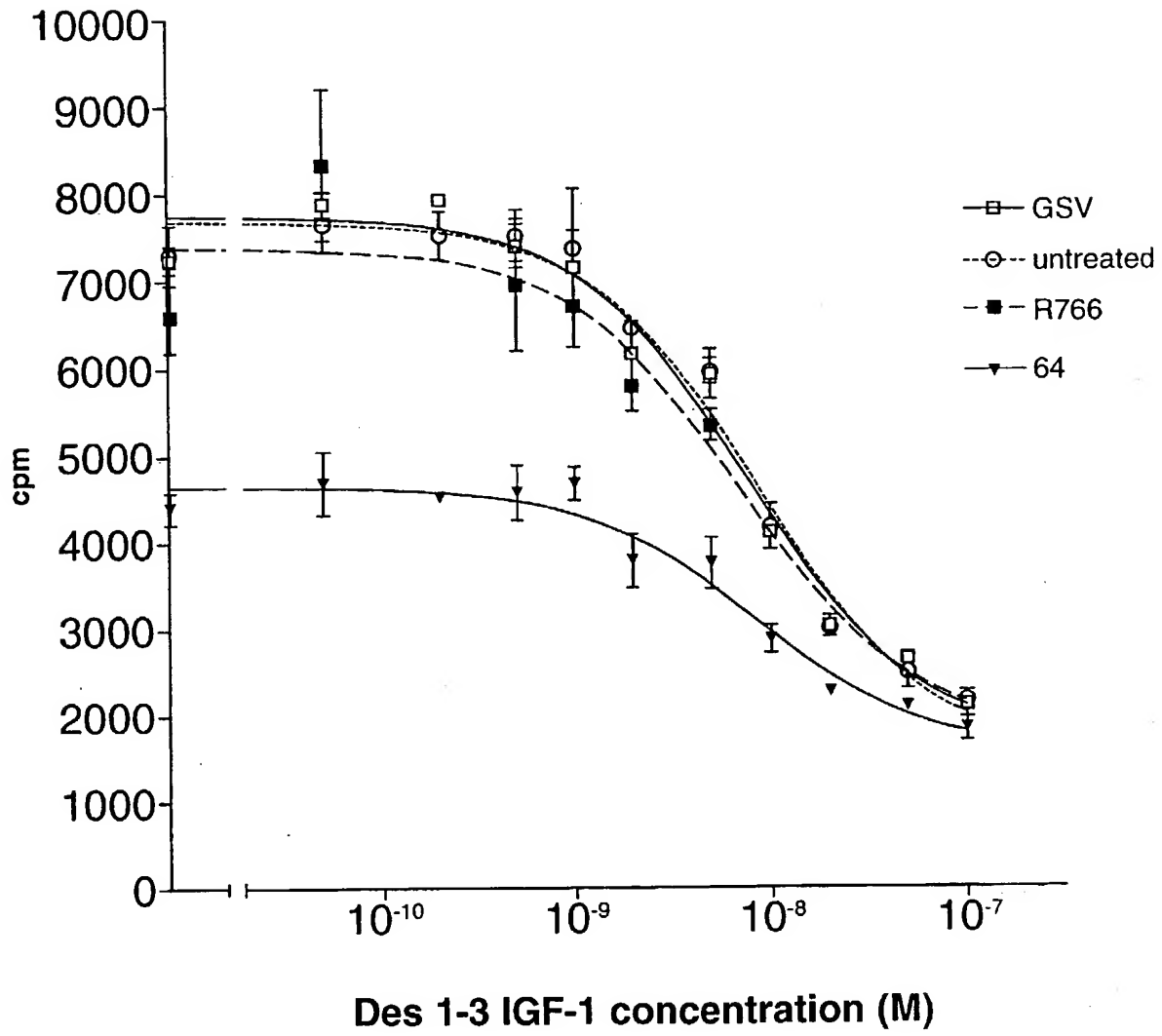


Figure 14

Substitute Sheet
(Rule 26) RO/AU

Effect of antisense oligonucleotides on IGF-1 receptor levels on the surface of keratinocytes:
Competition Assay - ^{125}I IGF-1 vs Des 1-3 IGF-1

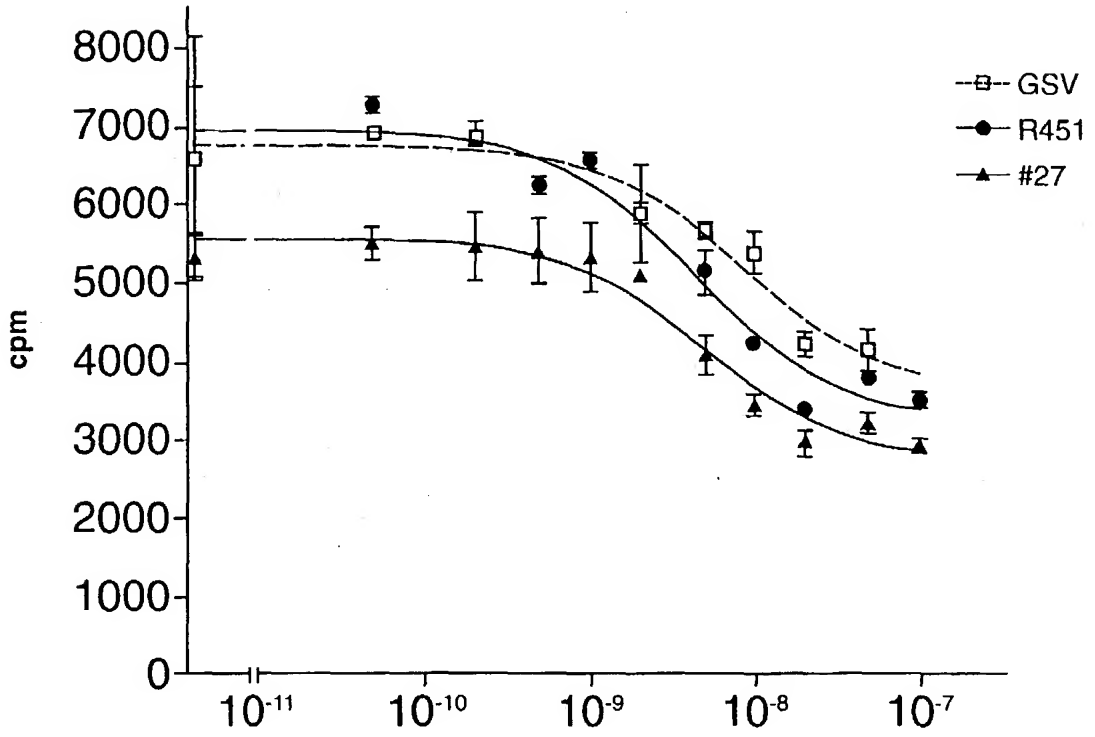


Figure 15

H&E stained sections of (A) psoriatic skin biopsy prior to grafting and (B) 49 day old psoriatic skin graft using skin from same donor

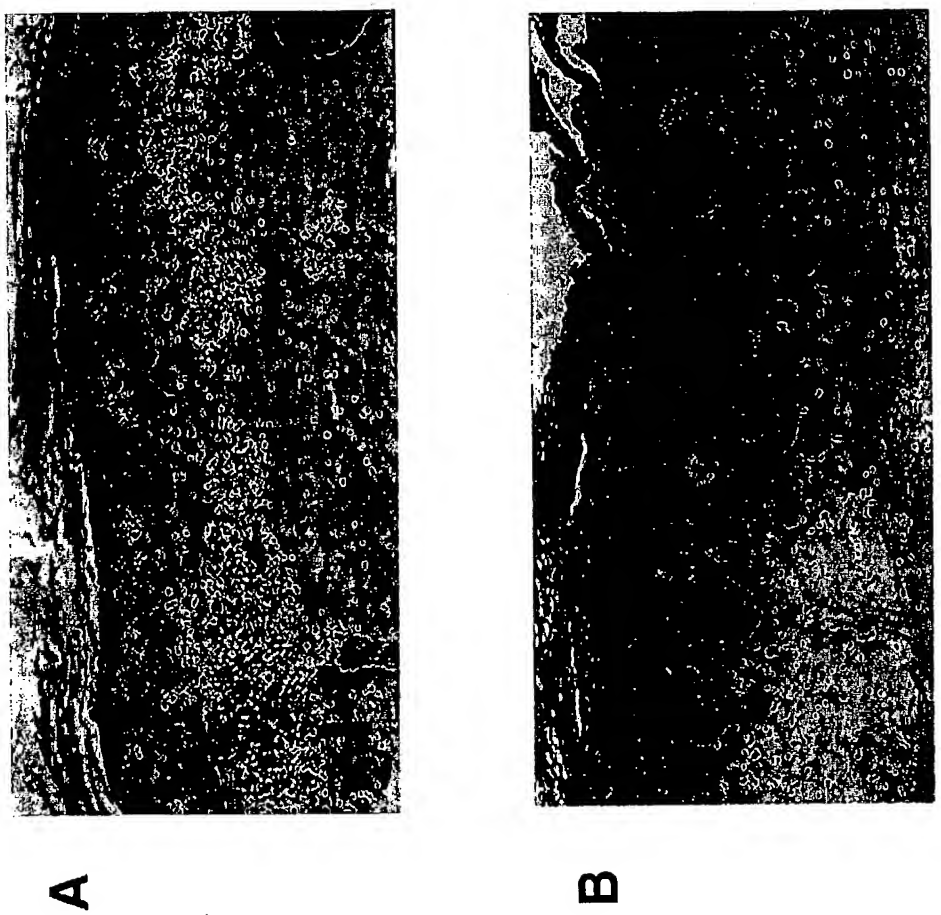


Figure 16

Uptake of oligonucleotide after intradermal injection into psoriatic skin graft on a nude mouse

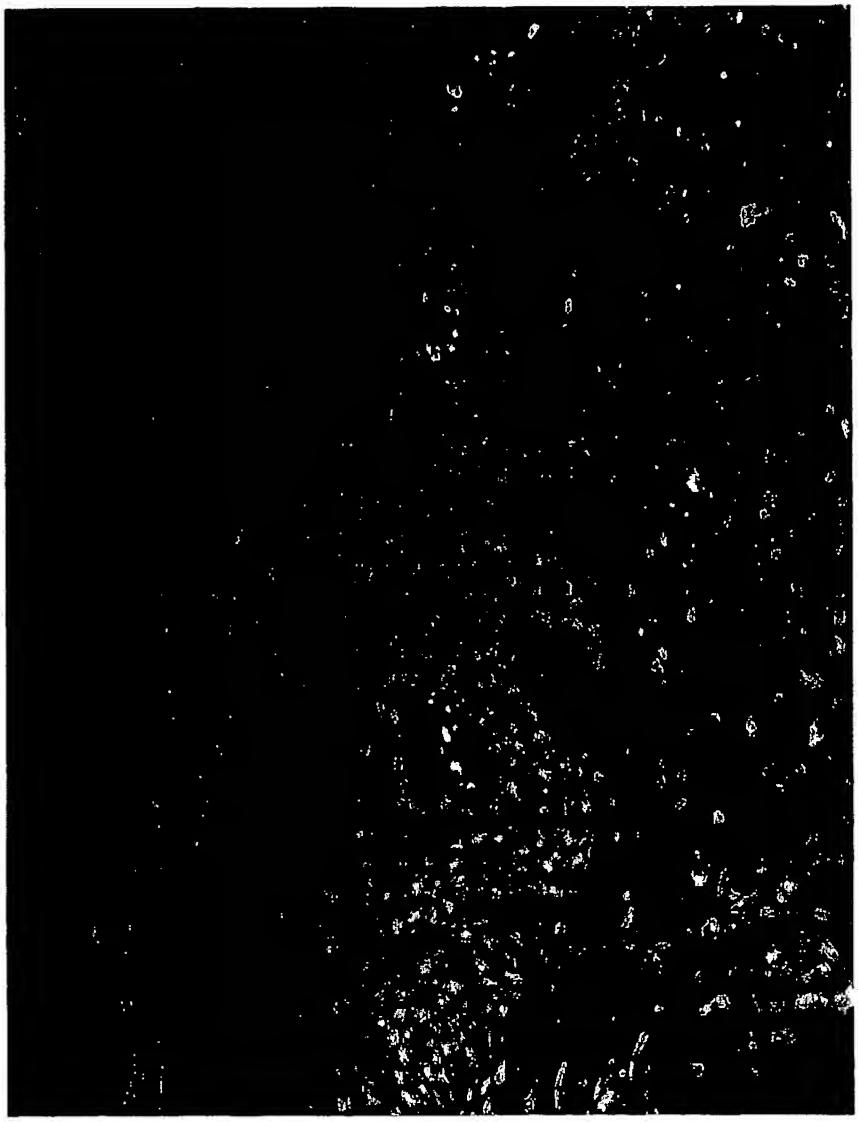


Figure 17

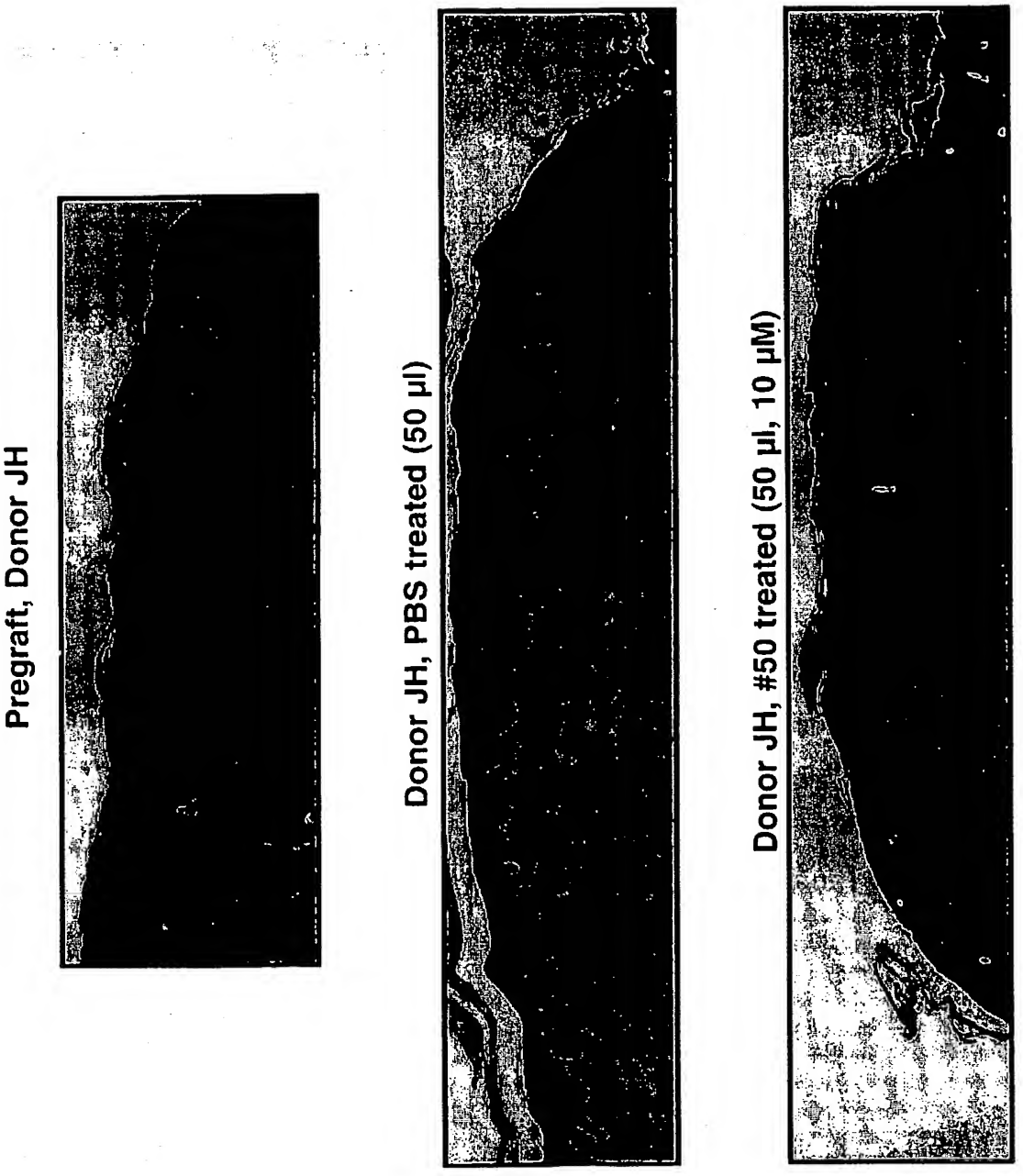

Pregraft, Donor JH
Donor JH, PBS treated (50 µl)
Donor JH, #50 treated (50 µl, 10 µM)

Figure 18a

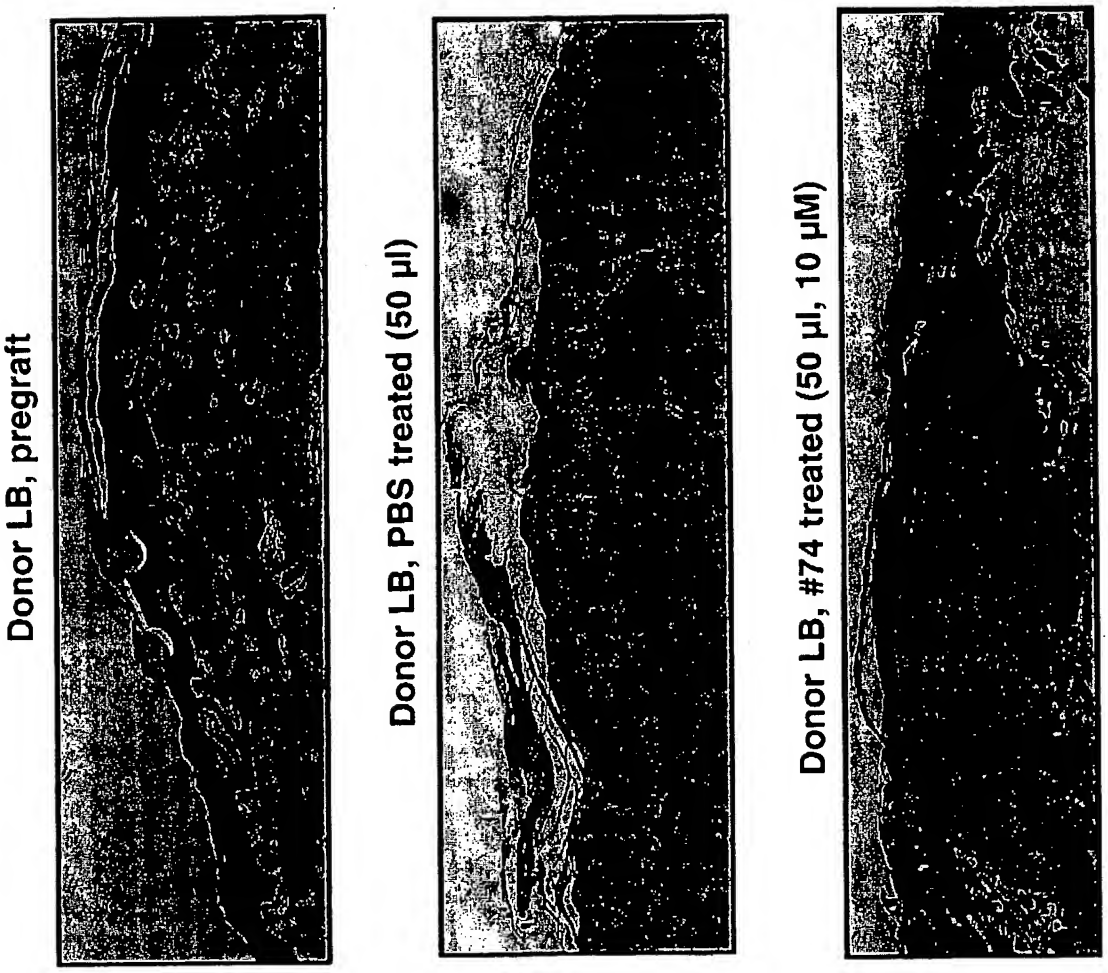

Donor LB, pregraft
Donor LB, PBS treated (50 µl)
Donor LB, #74 treated (50 µl, 10 µM)

Figure 18b

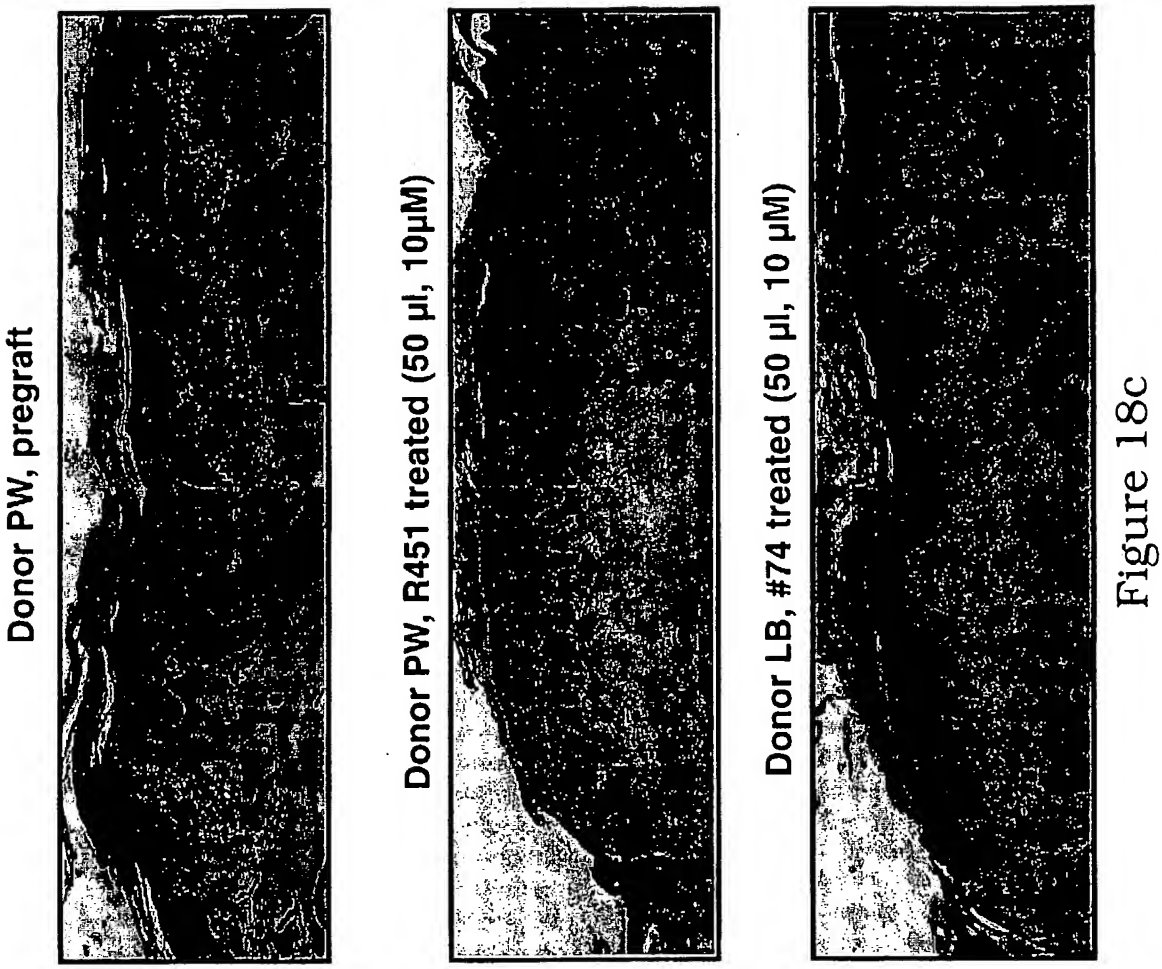

Donor PW, pregraft
Donor PW, R451 treated (50 µl, 10µM)
Donor LB, #74 treated (50 µl, 10 µM)

Figure 18c

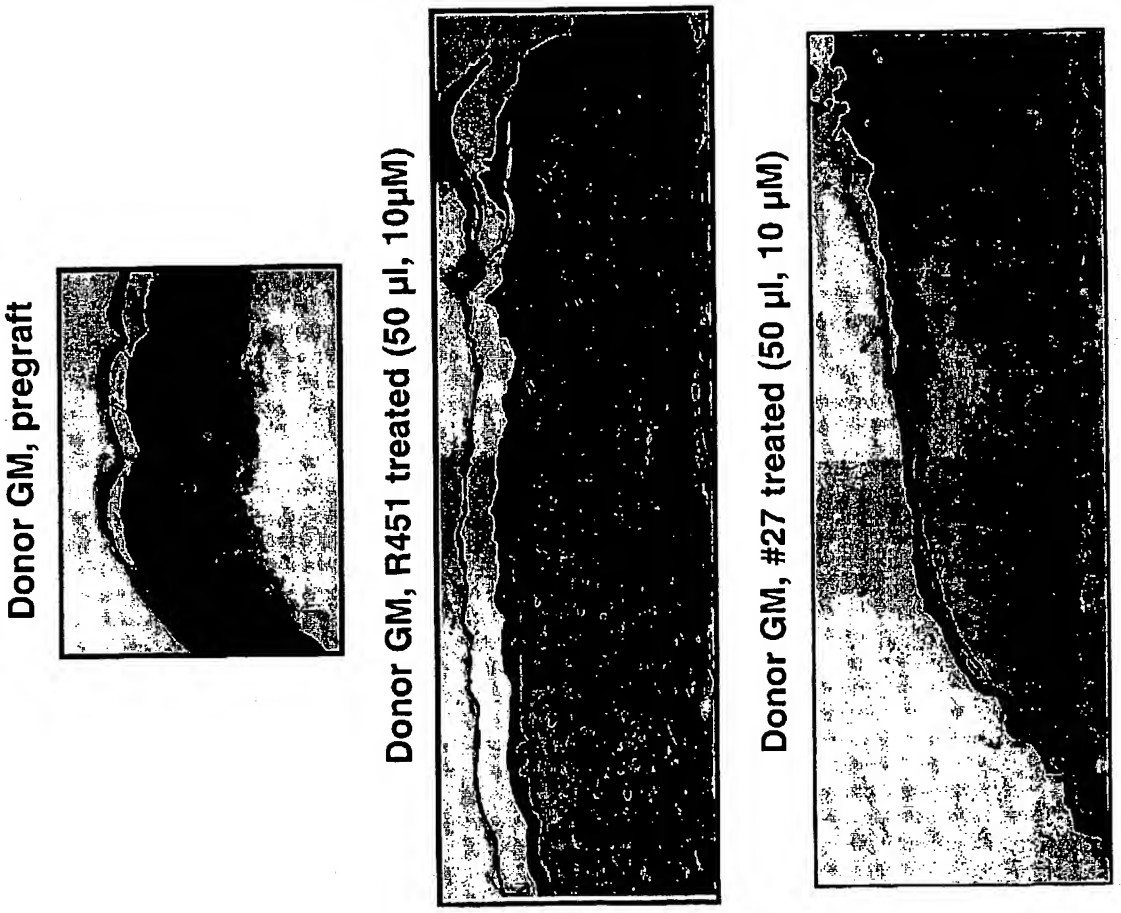
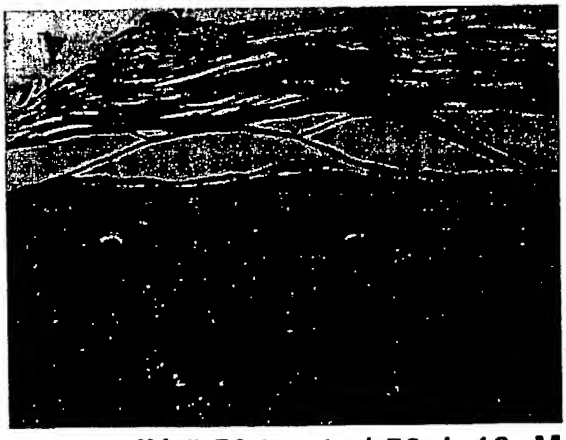

Donor GM, pregraft
Donor GM, R451 treated (50 µl, 10µM)
Donor GM, #27 treated (50 µl, 10 µM)

Figure 18d

Donor JH Pregraft

Donor JH PBS treated 50ul

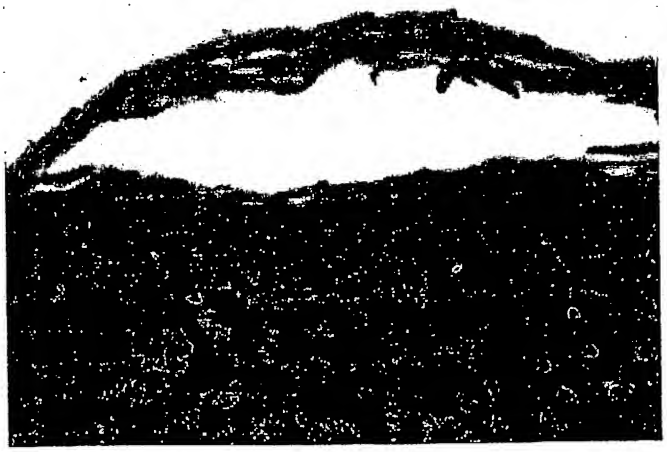


Donor JH # 50 treated 50ul, 10uM

Figure 19a

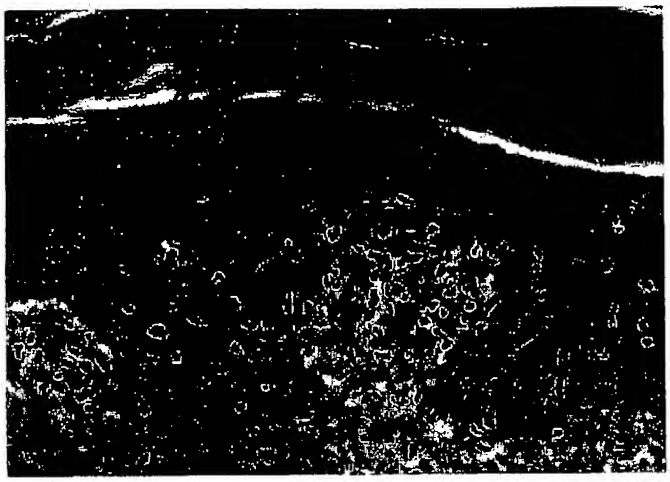
Donor LB Pregraft

Donor LB PBS treated 50ul



Donor LB # 74 treated 50ul, 10uM

Figure 19b



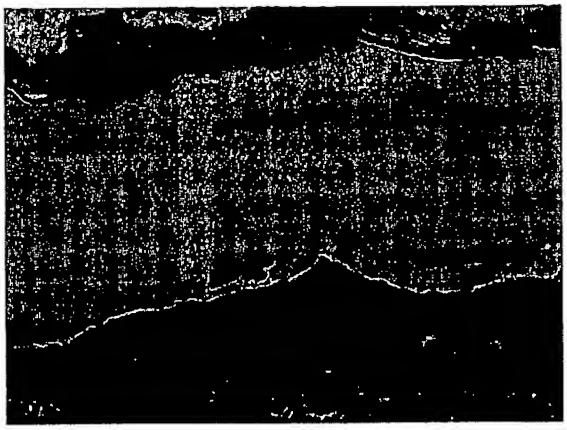
Donor PW Pregraft

Donor PW R451 treated 50ul, 10um

Donor PW # 74 treated 50ul, 10uM

Figure 19c

Donor GM Pregraft



Donor GM R451 treated 50ul, 10um

Donor GM # 27 treated 50ul, 10uM

Figure 19d

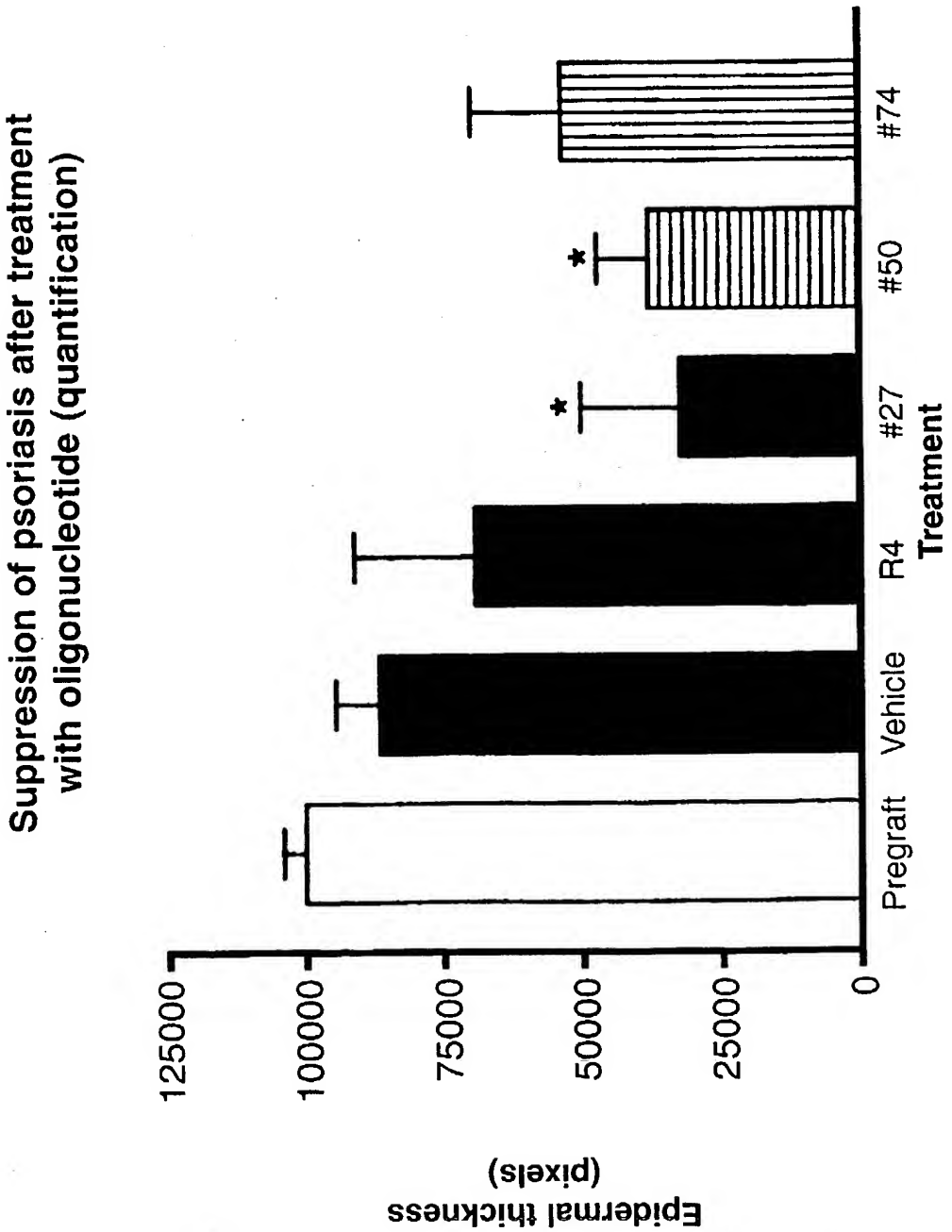

Suppression of psoriasis after treatment with oligonucleotide (quantification)
Epidermal thickness (pixels)
125000
100000
75000
50000
25000
0
Pregraft
Vehicle
R4
#27
#50
#74
Treatment
*
*

Figure 20

αhKi-67

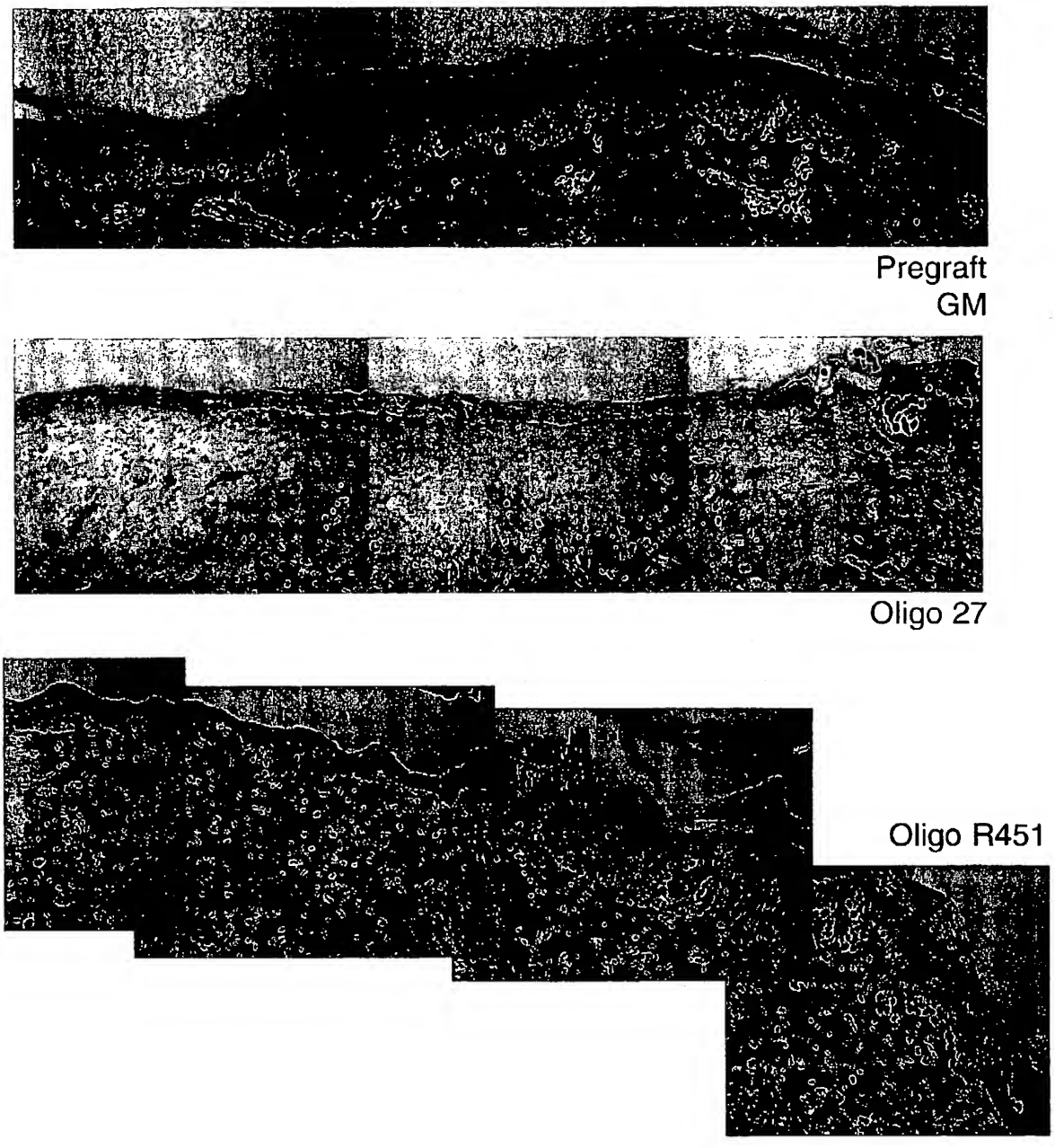


Figure 21

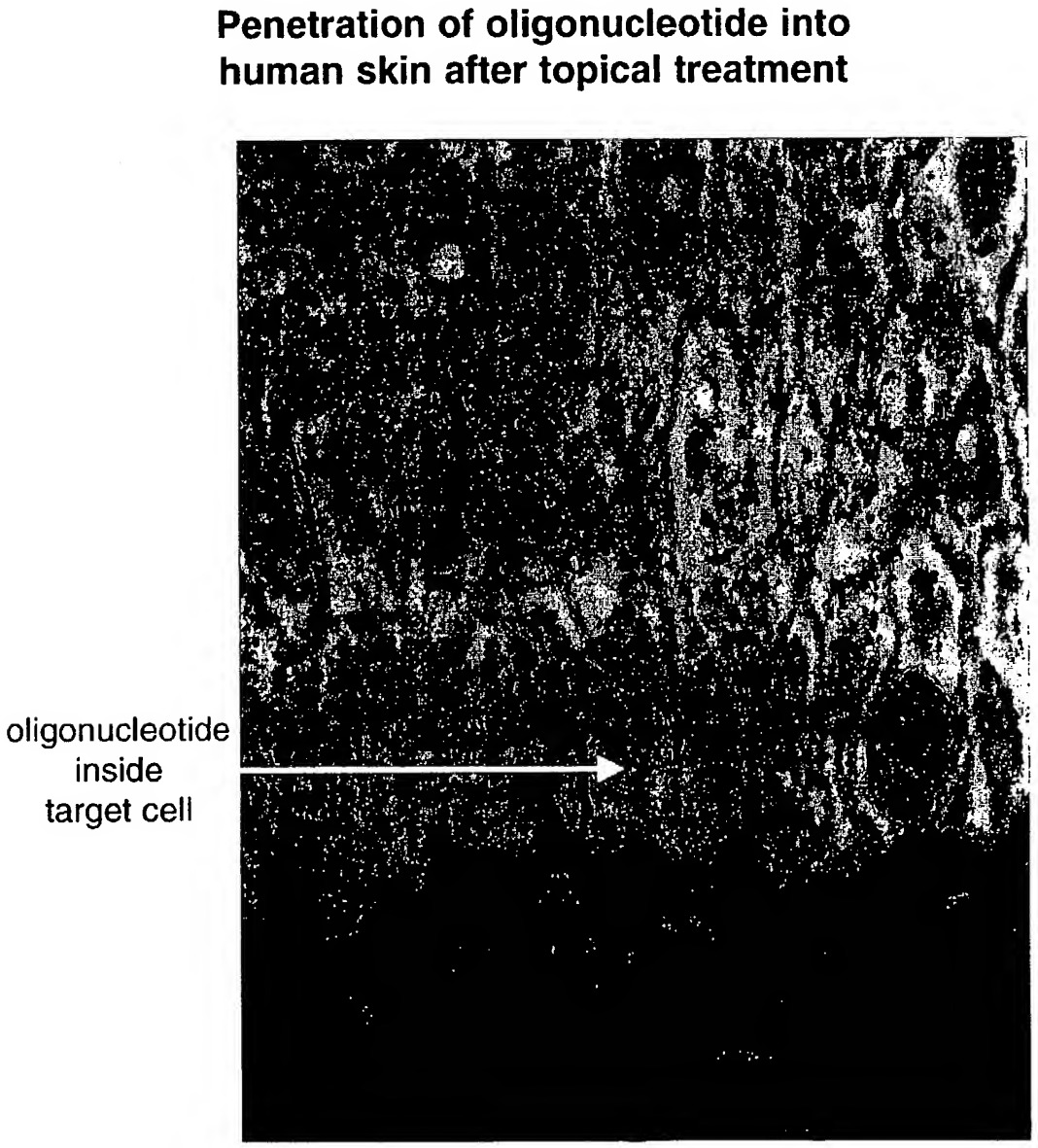

Penetration of oligonucleotide into human skin after topical treatment
oligonucleotide inside target cell

Figure 22

Penetration of oligonucleotide into human skin after topical gel formulation

Figure 23

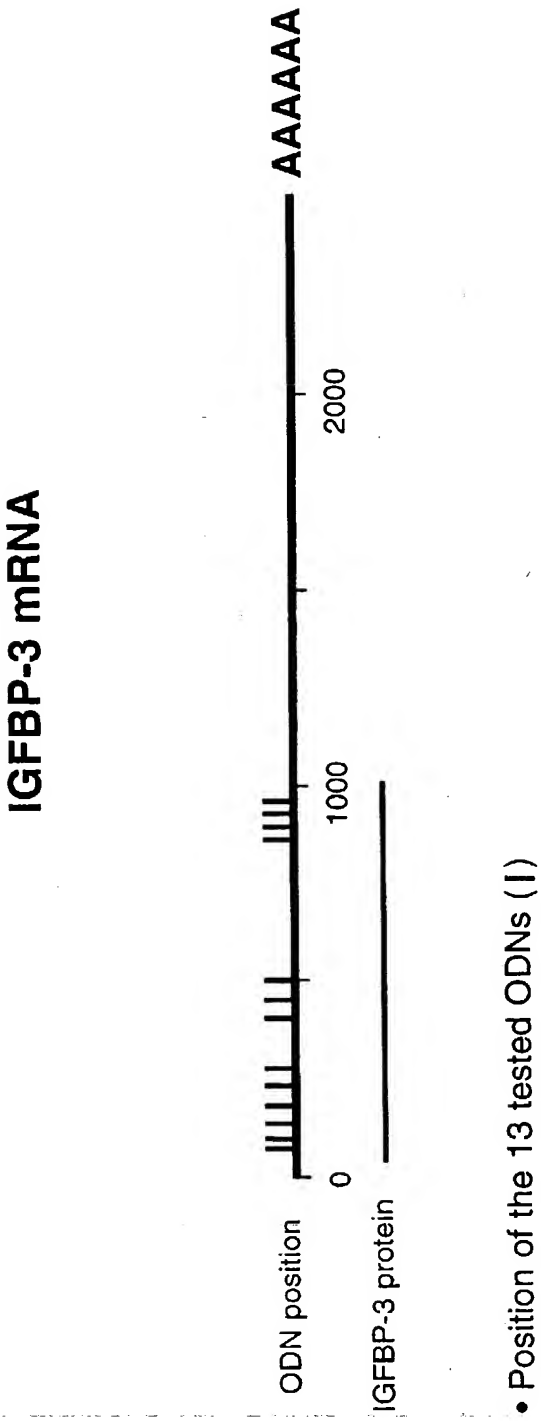

IGFBP-3 mRNA
AAAAAA
0
1000
2000
ODN position
IGFBP-3 protein
• Position of the 13 tested ODNs (I)
• mRNA transcript length = 2.5 Kb
• coding sequence 133-1009

Figure 24

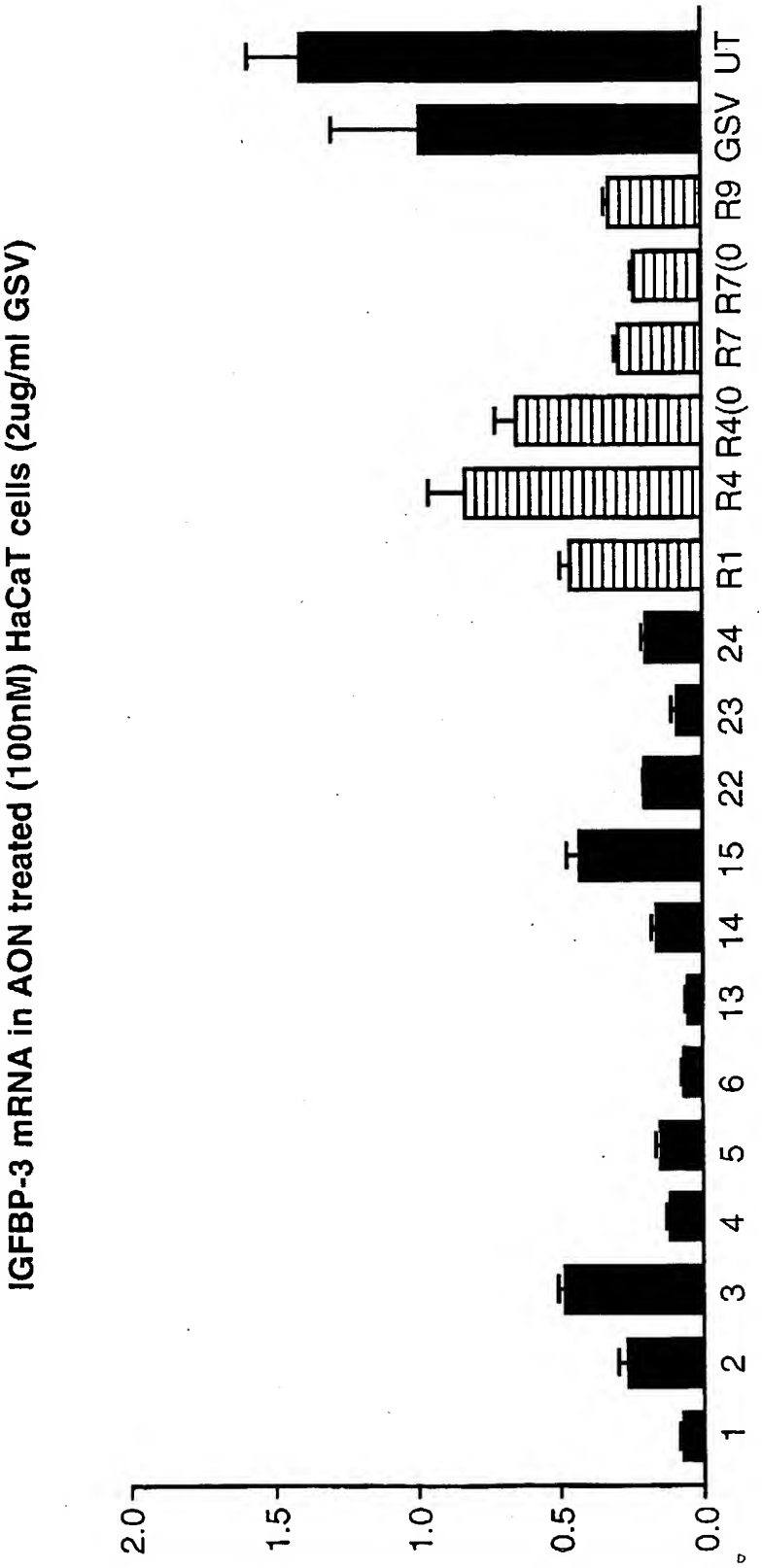

IGFBP-3 mRNA in AON treated (100nM) HaCaT cells (2ug/ml GSV)
relative IGFBP-3 mRNA
2.0
1.5
1.0
0.5
0.0
1
2
3
4
5
6
13
14
15
22
23
24
R1
R4
R4(0
R7
R7(0
R9
GSV
UT
Treatment

Figure 25a

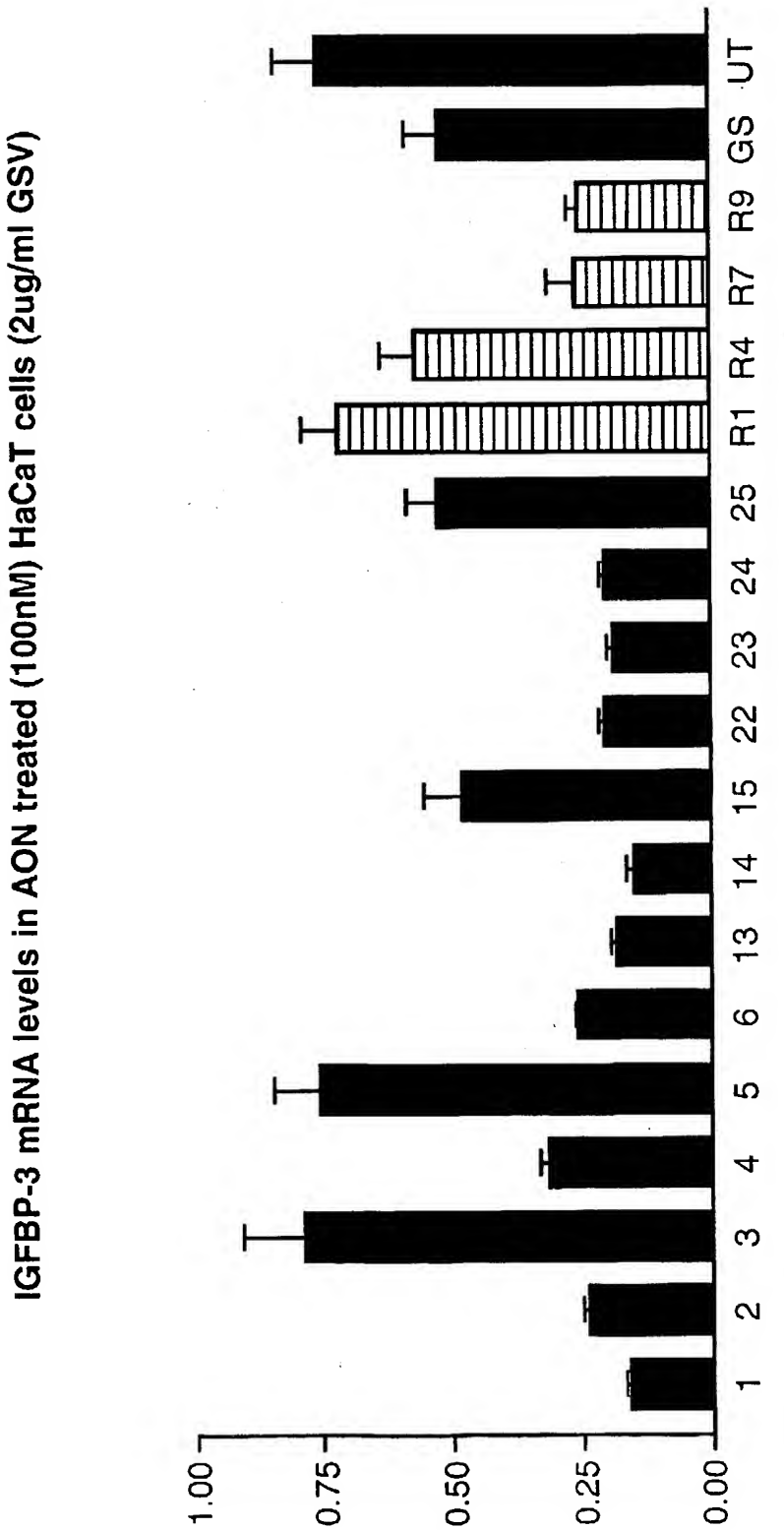

IGFBP-3 mRNA levels in AON treated (100nM) HaCaT cells (2ug/ml GSV)
relative IGFBP-3 mRNA
1.00
0.75
0.50
0.25
0.00
1
2
3
4
5
6
13
14
15
22
23
24
25
R1
R4
R7
R9
GS
UT
Treatment

Figure 25b

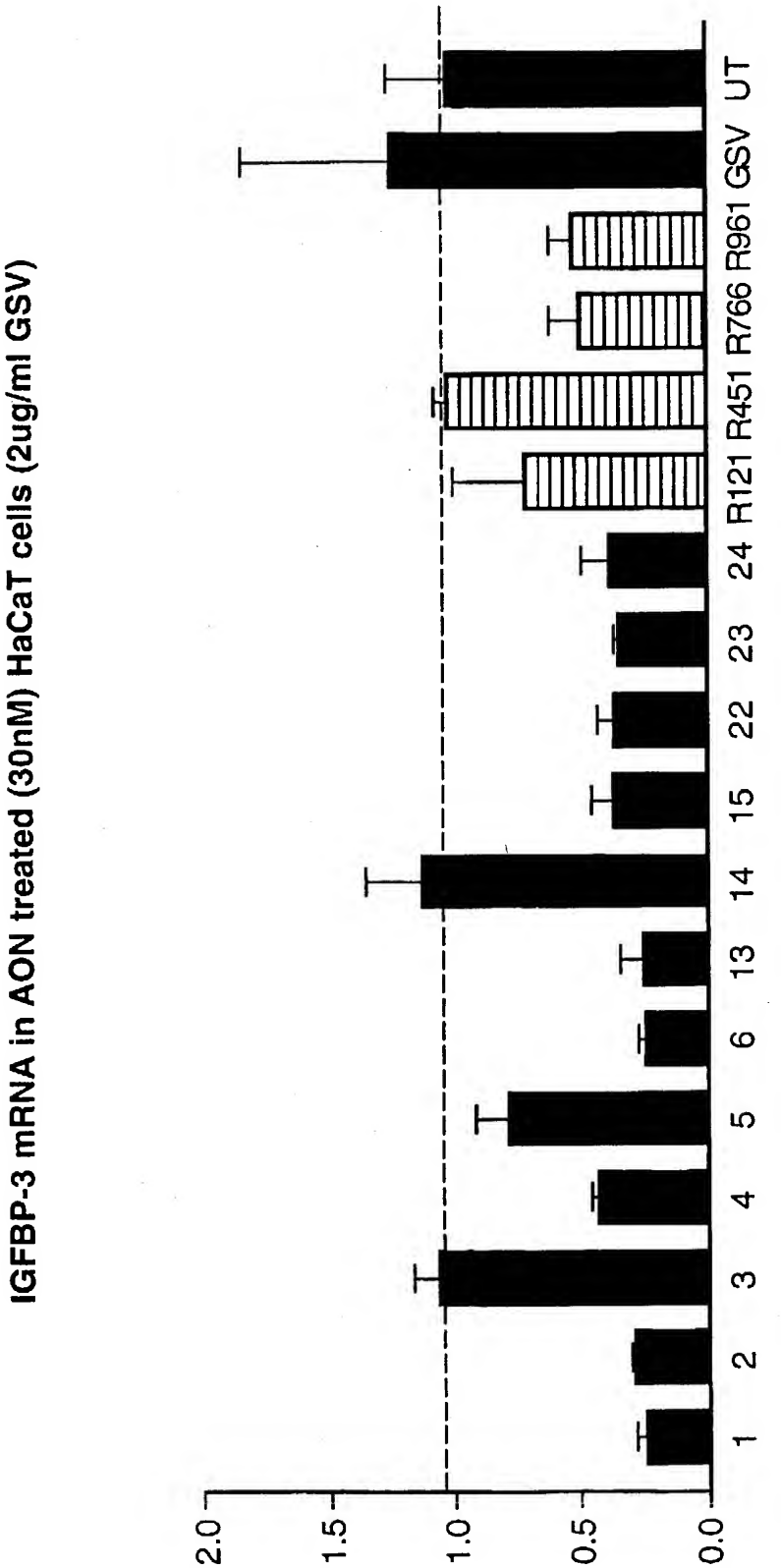

IGFBP-3 mRNA in AON treated (30nM) HaCaT cells (2ug/ml GSV)
relative IGFBP-3 mRNA
2.0
1.5
1.0
0.5
0.0
1
2
3
4
5
6
13
14
15
22
23
24
R121
R451
R766
R961
GSV
UT
Treatment

Figure 25c

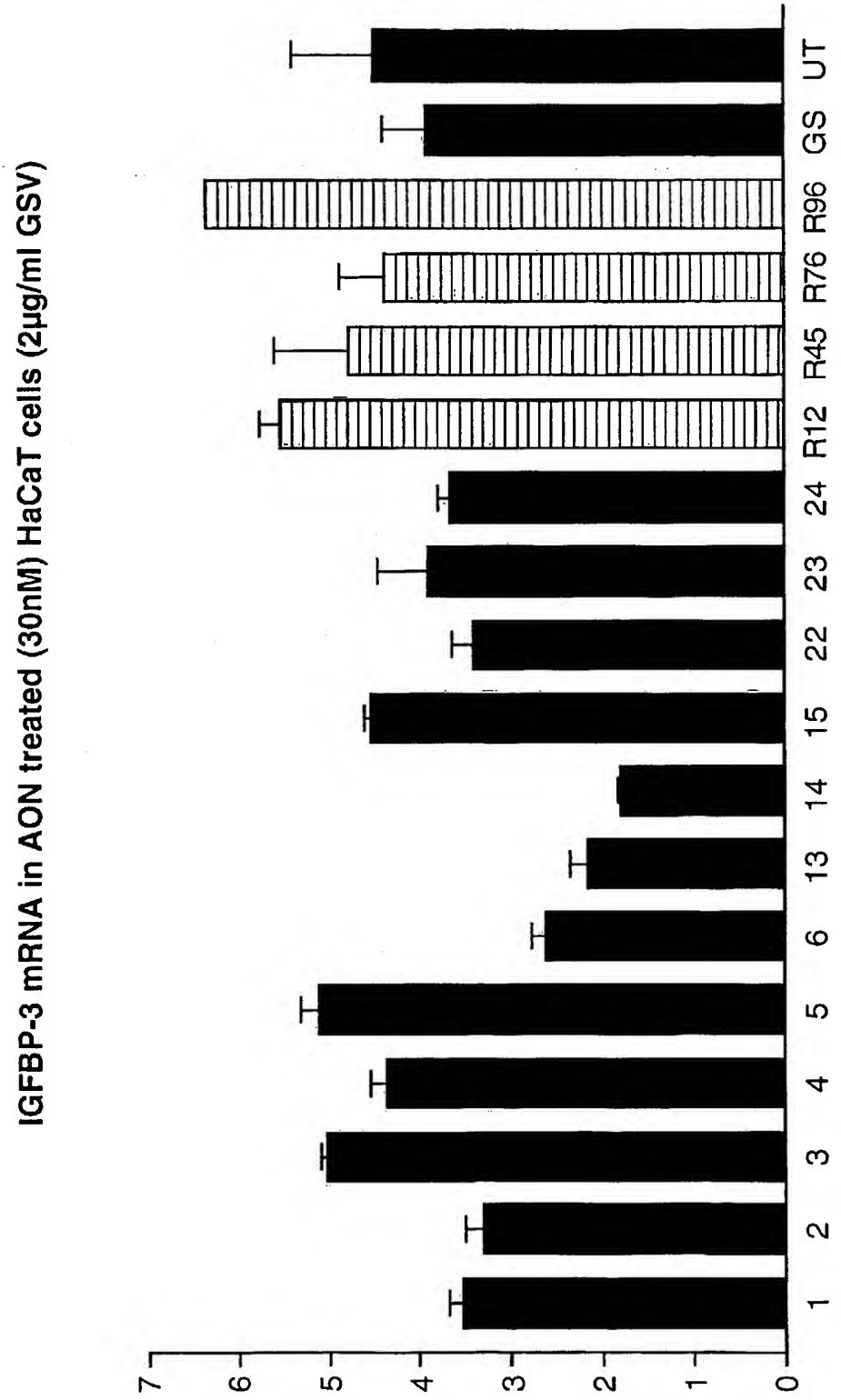

IGFBP-3 mRNA in AON treated (30nM) HaCaT cells (2µg/ml GSV)
relative IGFBP-3 mRNA
0
1
2
3
4
5
6
7
1
2
3
4
5
6
13
14
15
22
23
24
R12
R45
R76
R96
GS
UT
Treatment

Figure 25d

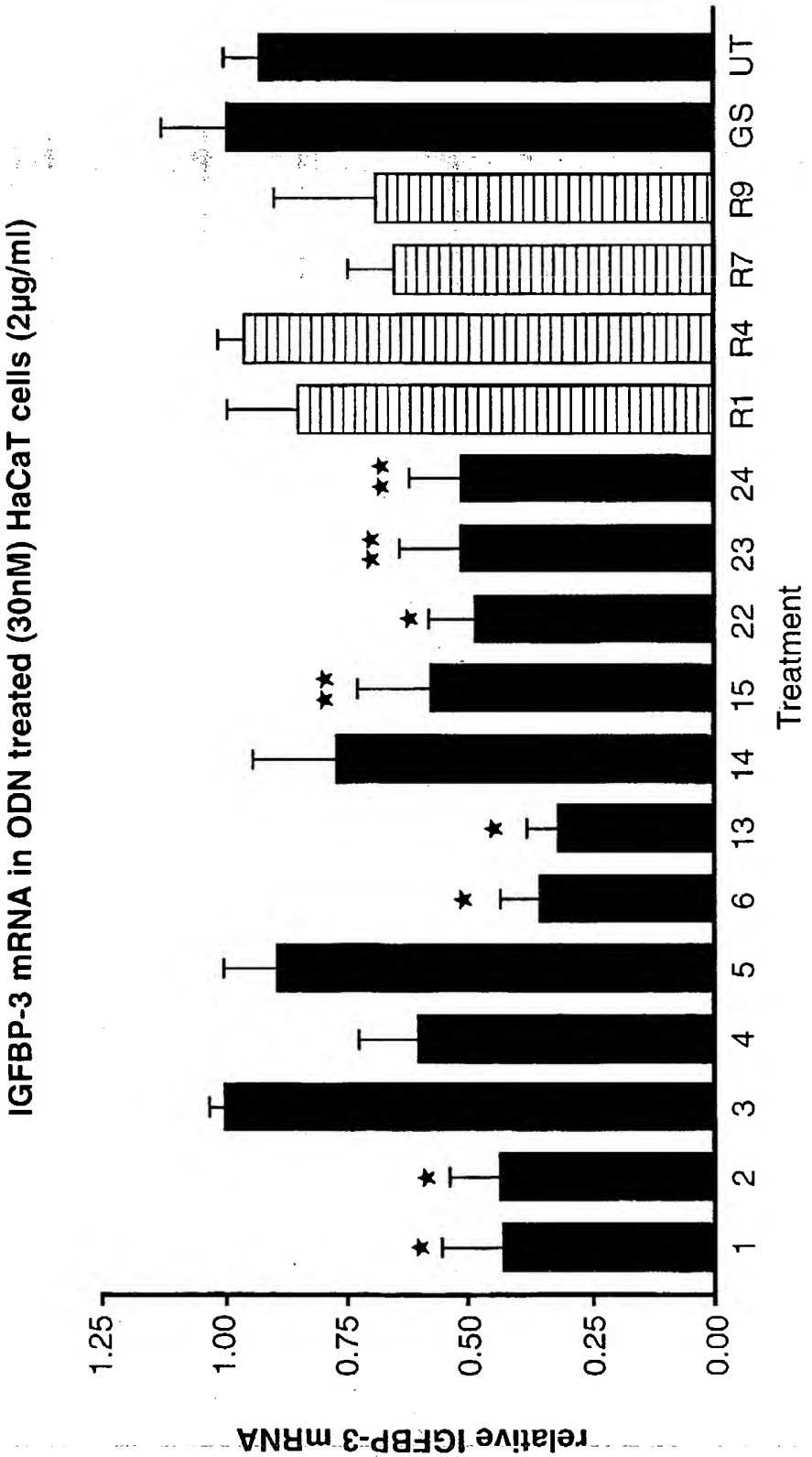

IGFBP-3 mRNA in ODN treated (30nM) HaCaT cells (2µg/ml)
relative IGFBP-3 mRNA
1.25
1.00
0.75
0.50
0.25
0.00
1
2
3
4
5
6
13
14
15
22
23
24
R1
R4
R7
R9
GS
UT
Treatment

Figure 26a

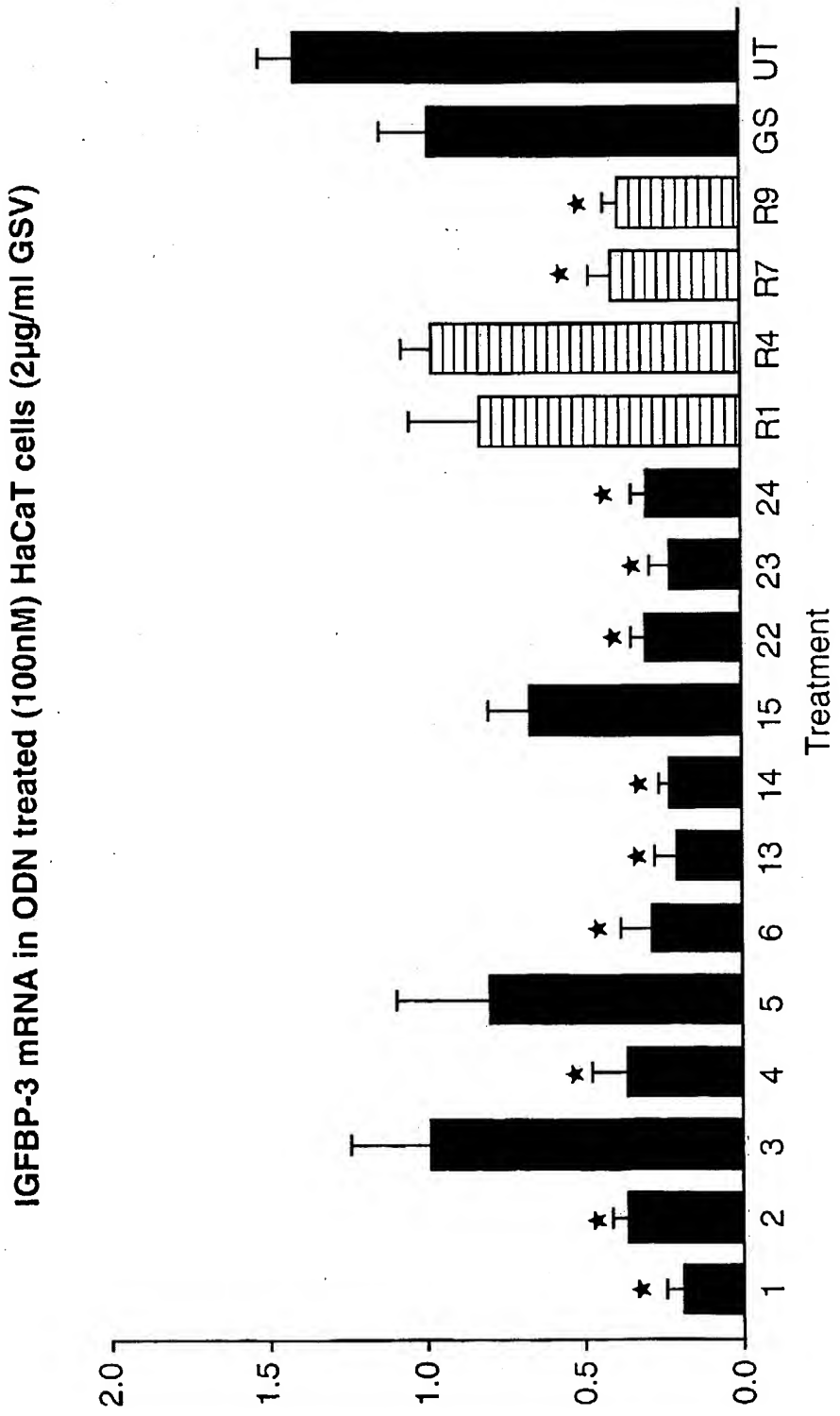

IGFBP-3 mRNA in ODN treated (100nM) HaCaT cells (2µg/ml GSV)
relative IGFBP-3 mRNA
2.0
1.5
1.0
0.5
0.0
1
2
3
4
5
6
13
14
15
22
23
24
R1
R4
R7
R9
GS
UT
Treatment

Figure 26b

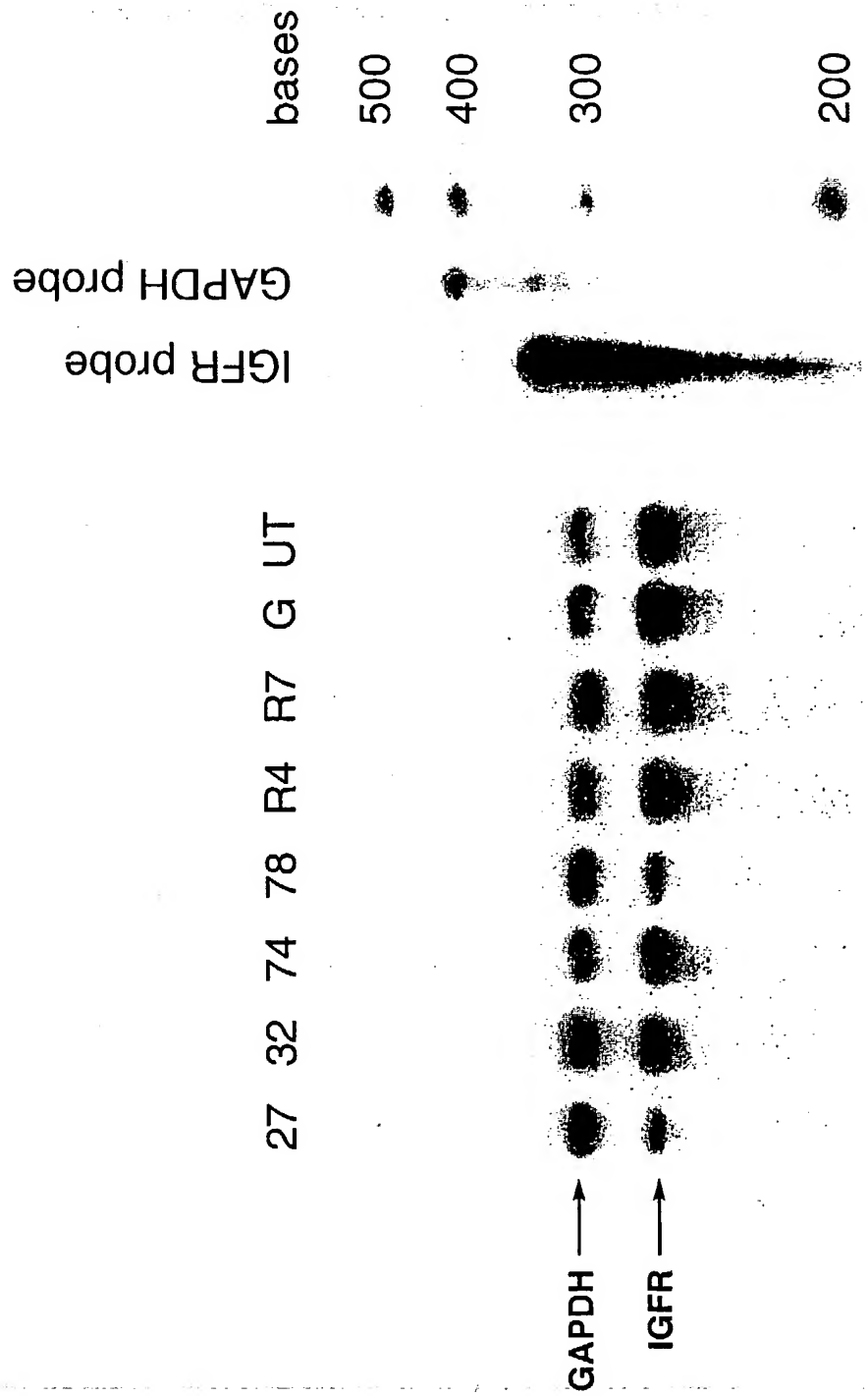

bases
500
400
300
200
GAPDH probe
IGFR probe
27
32
74
78
R4
R7
G
UT
GAPDH
IGFR

Figure 27a

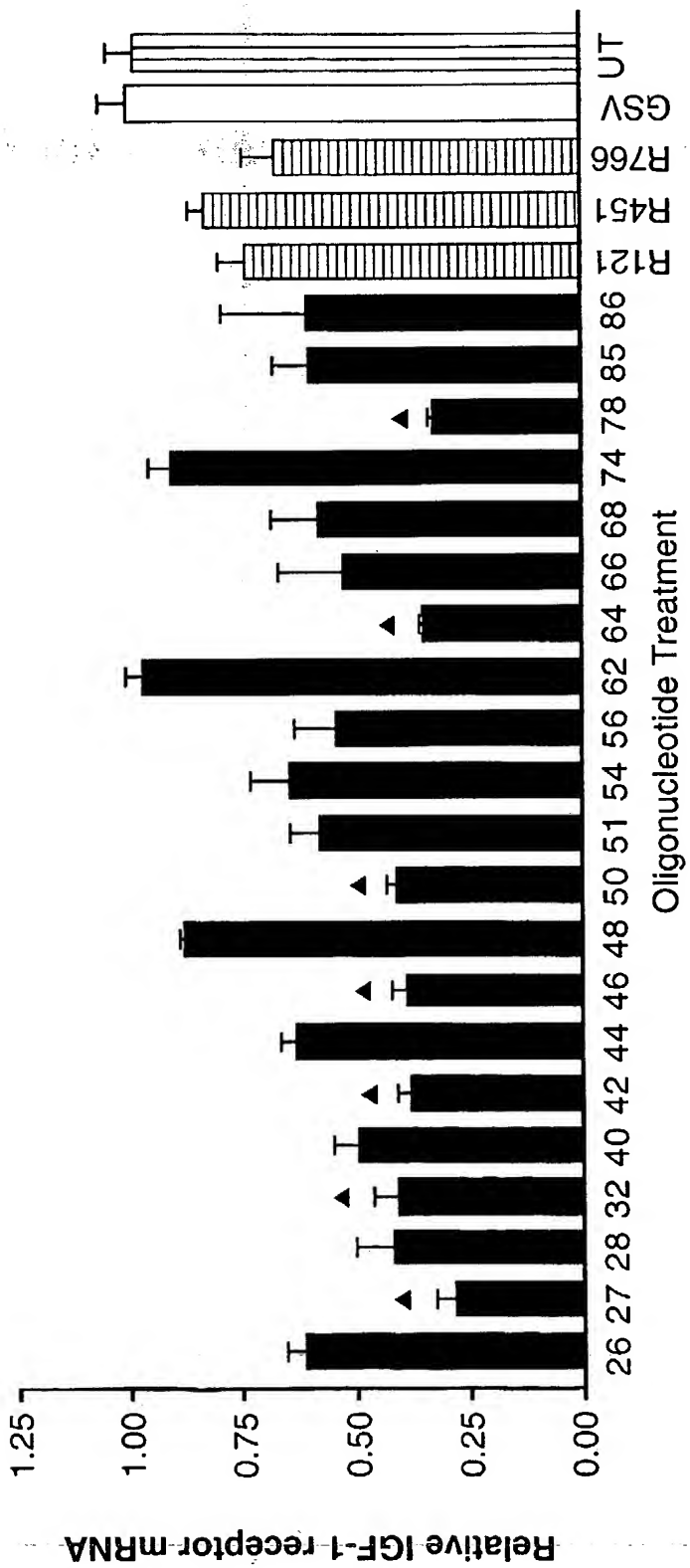

Relative IGF-1 receptor mRNA
0.00
0.25
0.50
0.75
1.00
1.25
26 27 28 32 40 42 44 46 48 50 51 54 56 62 64 66 68 74 78 85 86
R121
R451
R766
GSV
UT
Oligonucleotide Treatment

Figure 27 b

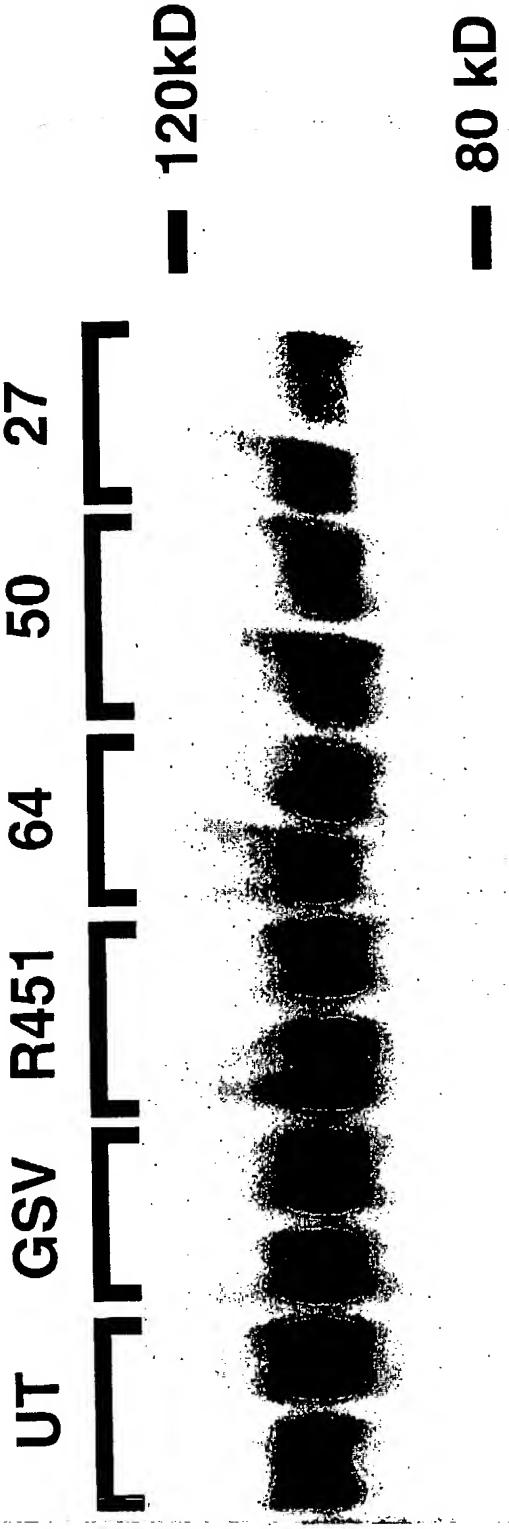


Figure 28a

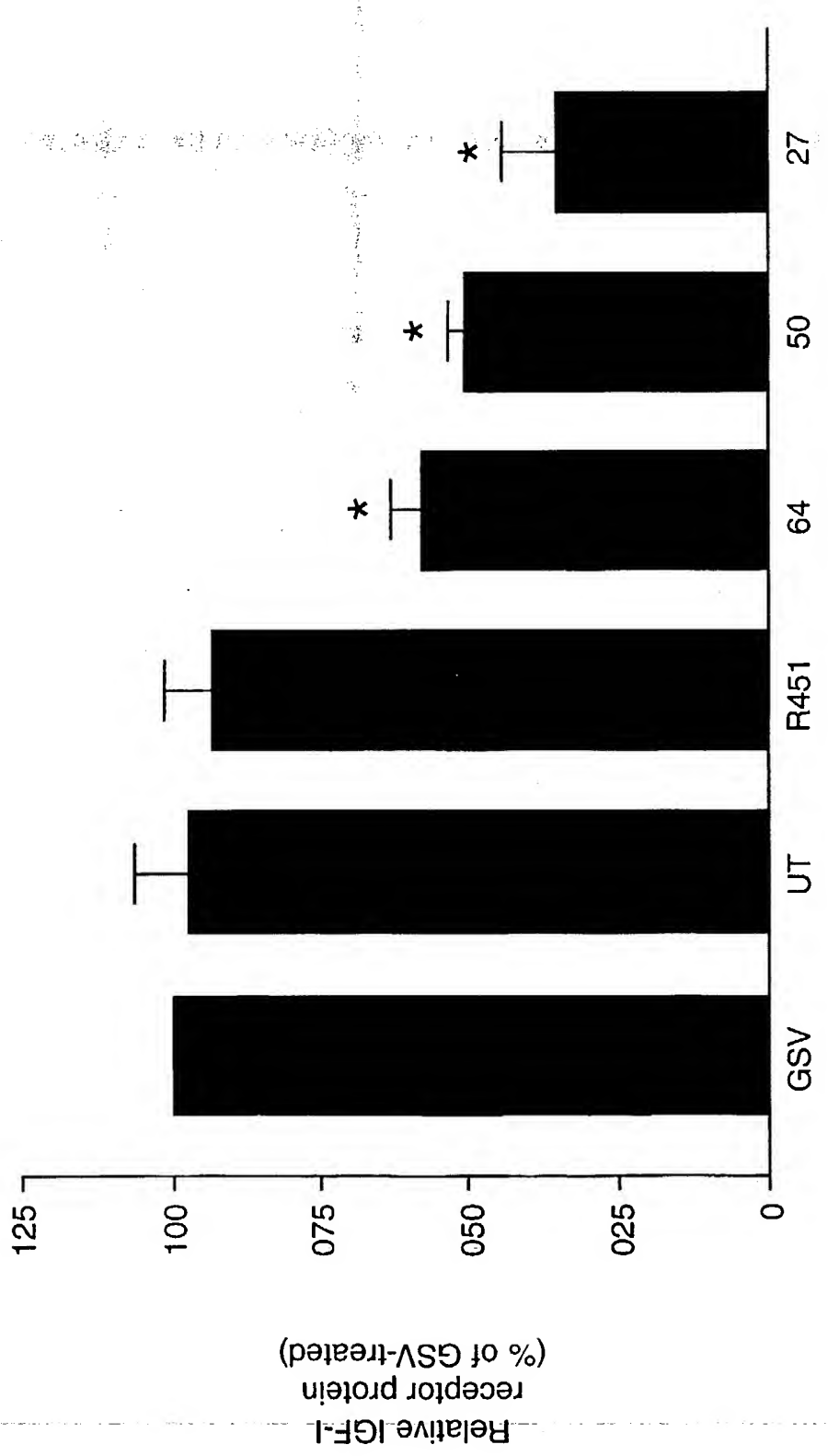

Relative IGF-I receptor protein (% of GSV-treated)
125
100
075
050
025
0
GSV
UT
R451
64
50
27
*
*
*

Figure 28b

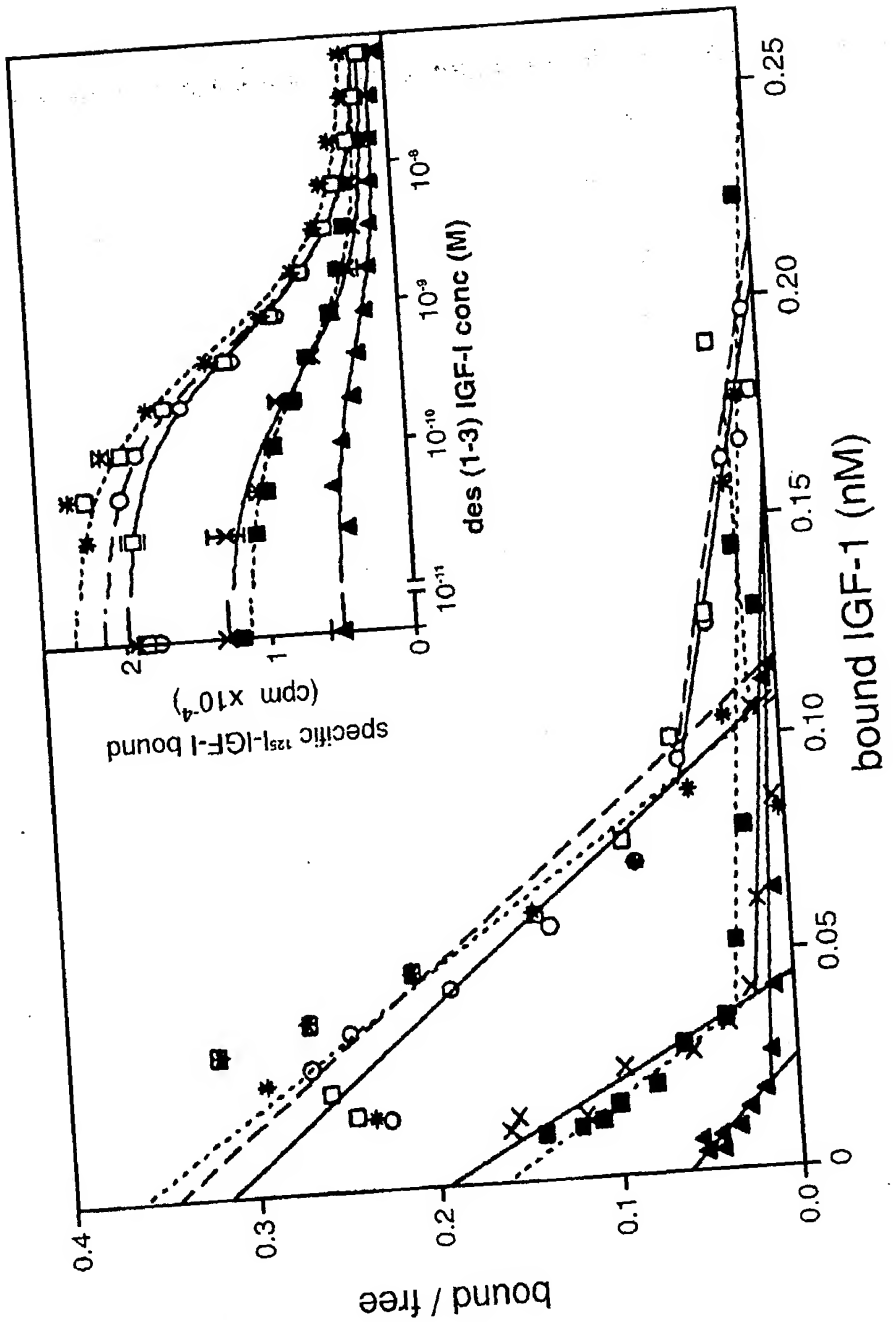

bound / free
0.4
0.3
0.2
0.1
0.0
bound IGF-1 (nM)
0
0.05
0.10
0.15
0.20
0.25
des (1-3) IGF-I conc (M)
10^{-8}
10^{-9}
10^{-10}
10^{-11}
0
1
2
specific ^{125}I-IGF-I bound
(cpm x10^{-4})

Figure 29

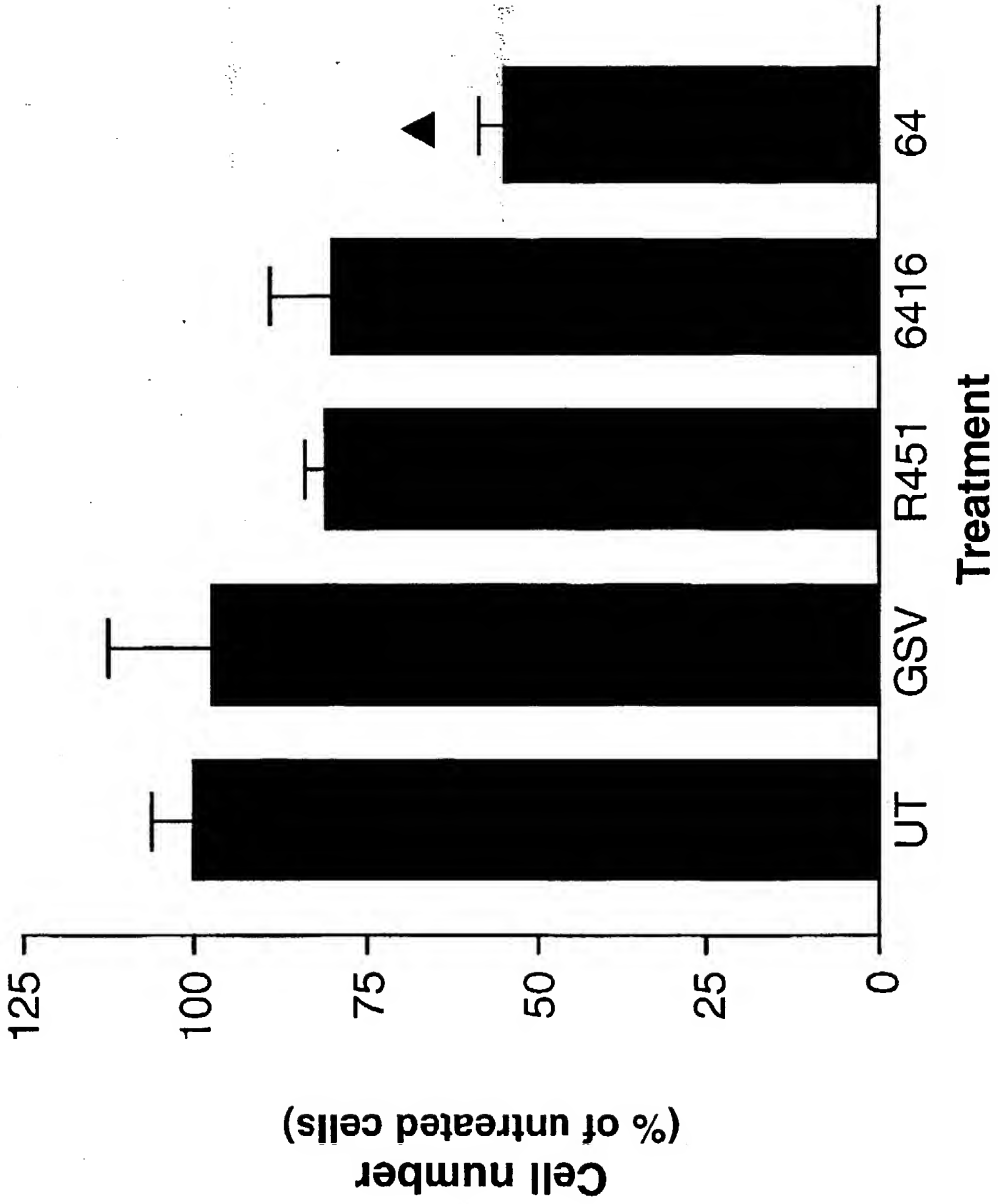

Cell number
(% of untreated cells)
125
100
75
50
25
0
UT
GSV
R451
6416
64
Treatment

Figure 30

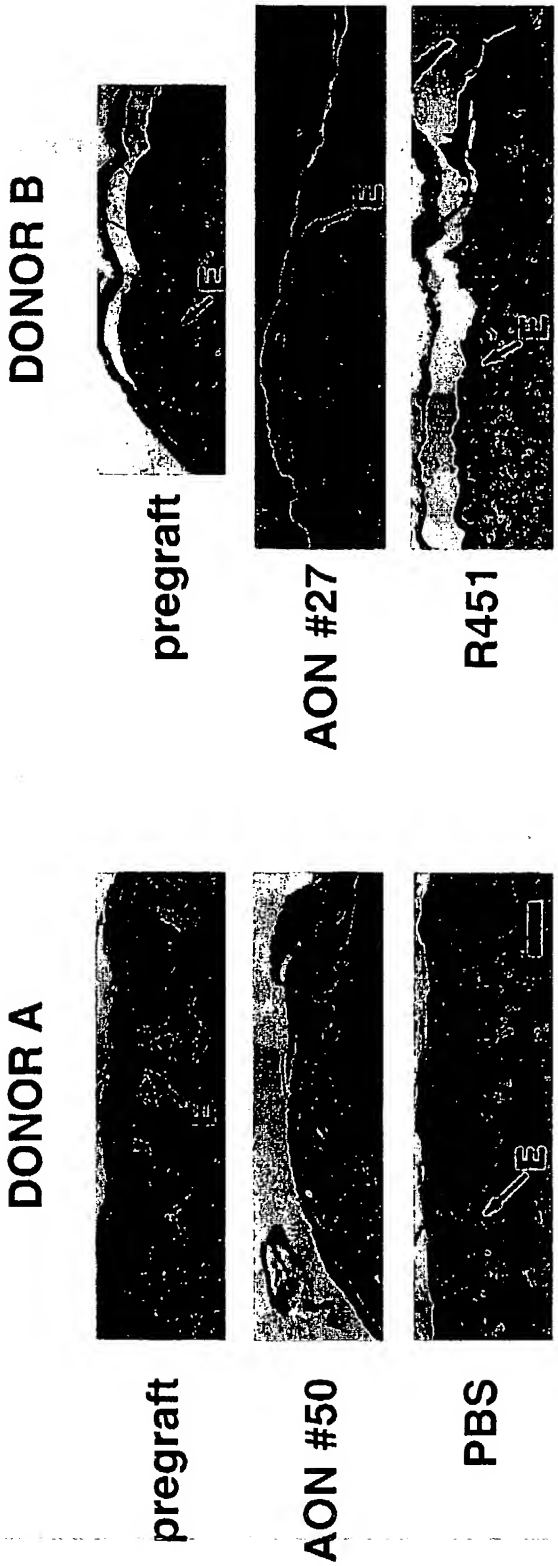


Figure 31a

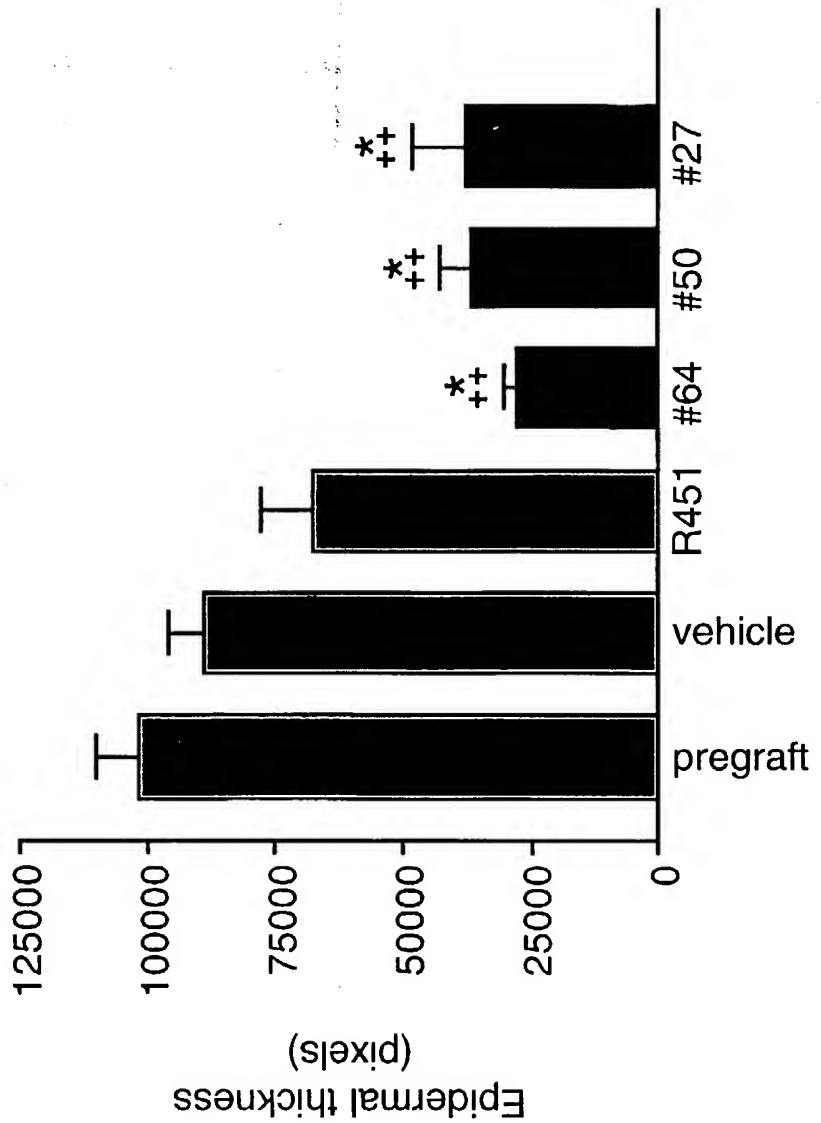

Epidermal thickness
(pixels)
125000
100000
75000
50000
25000
0
pregraft
vehicle
R451
#64
#50
#27
*++
*++
*++
*++

Figure 31b

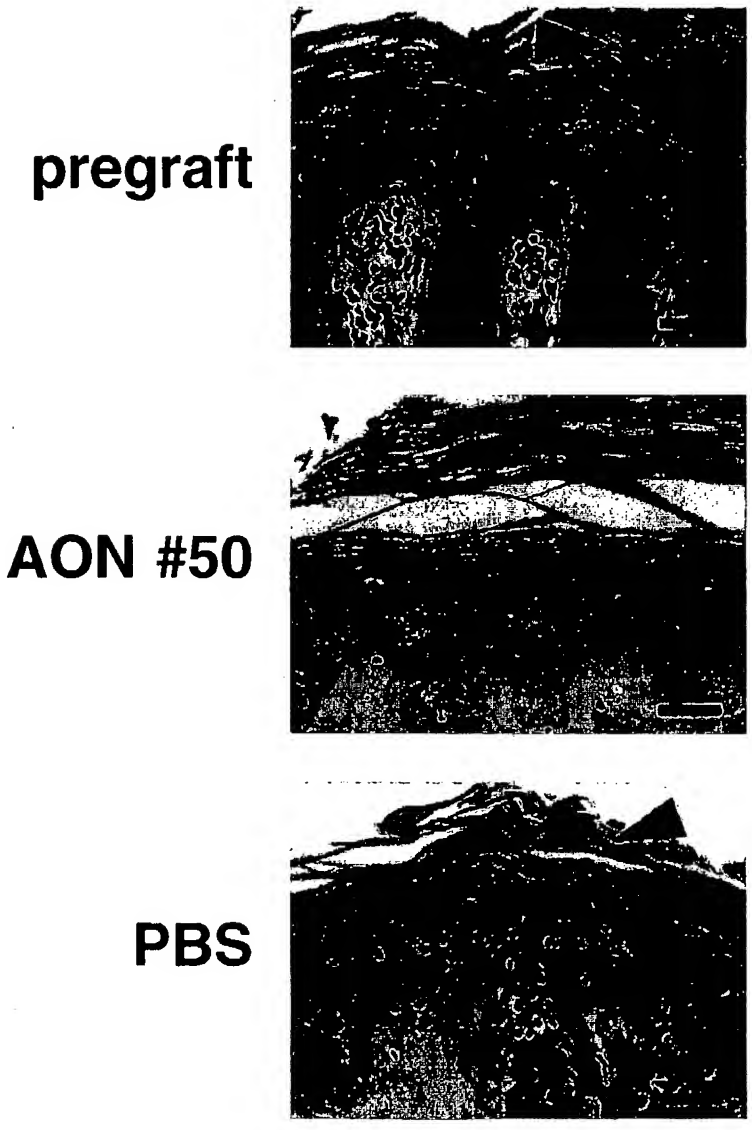


Figure 31c

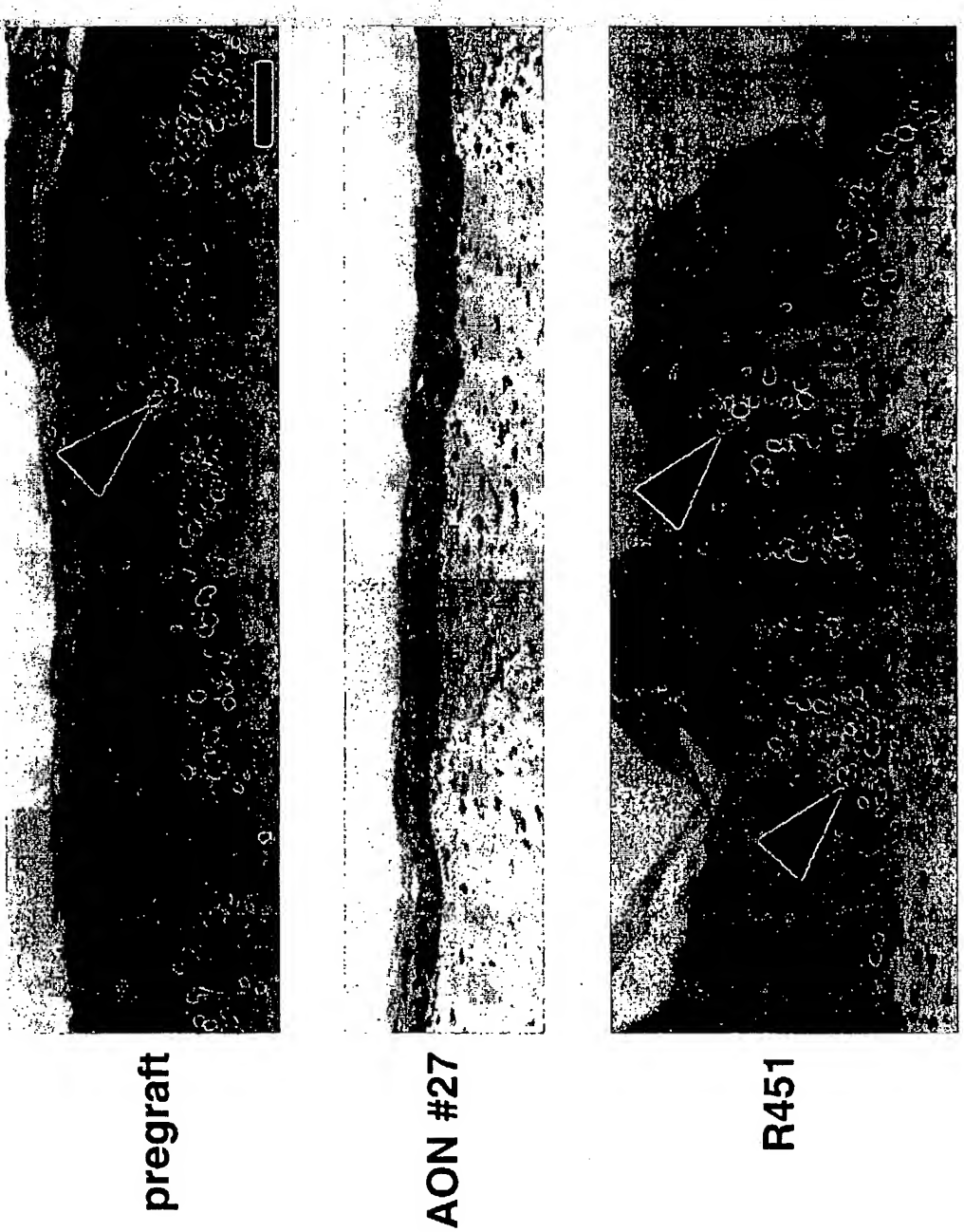


Figure 32a

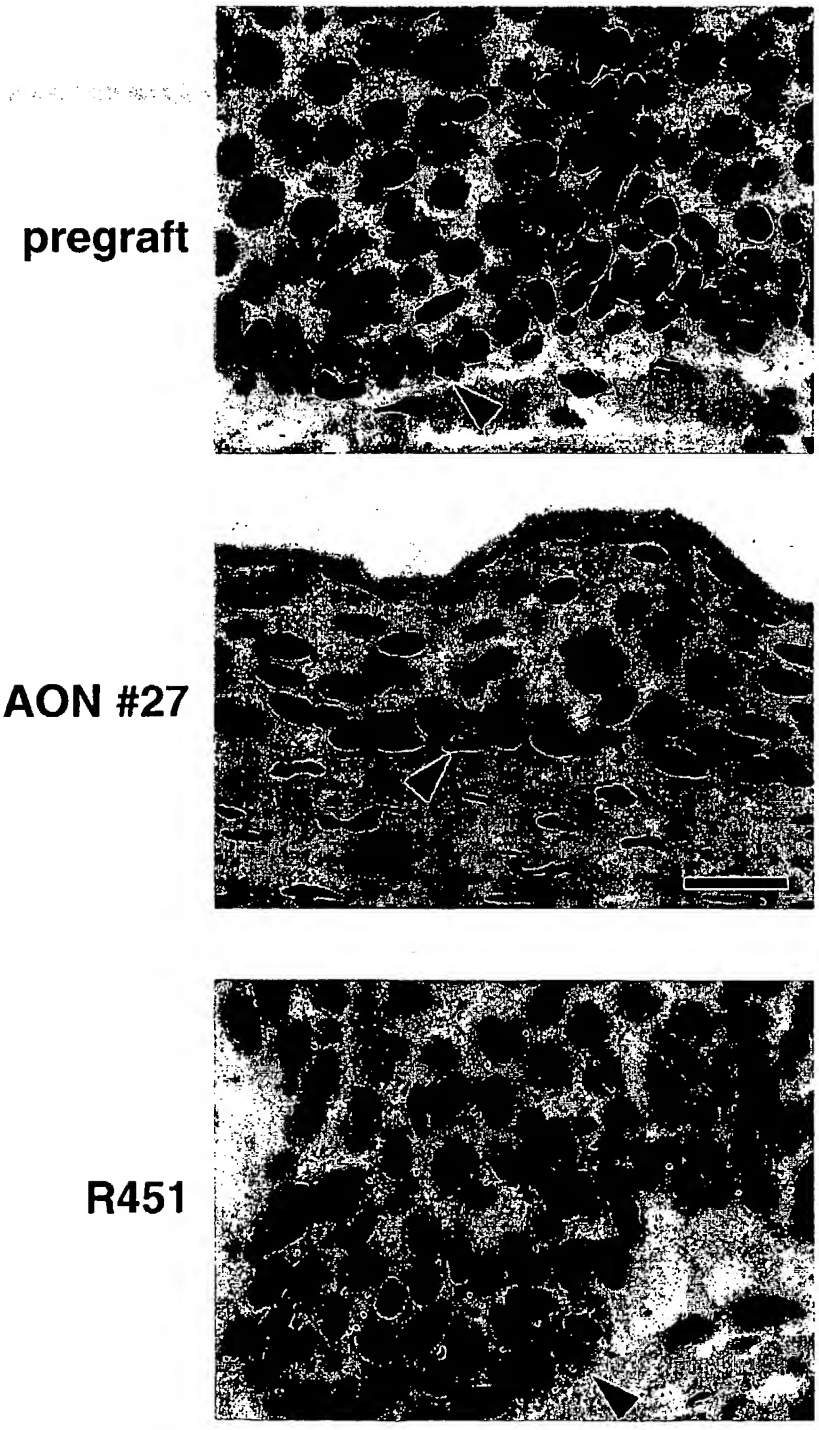


Figure 32b

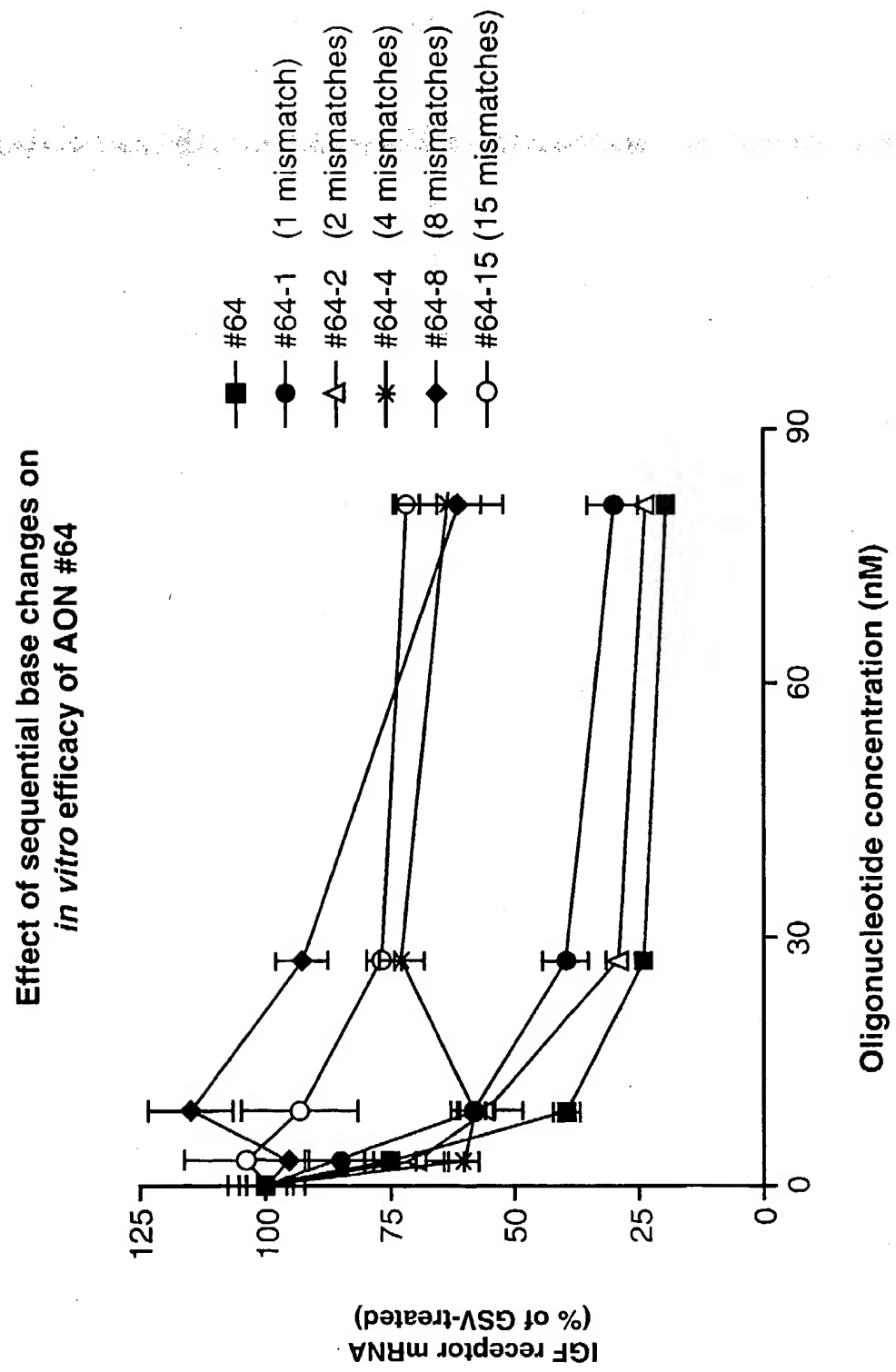

Effect of sequential base changes on
in vitro efficacy of AON #64
#64
#64-1 (1 mismatch)
#64-2 (2 mismatches)
#64-4 (4 mismatches)
#64-8 (8 mismatches)
#64-15 (15 mismatches)
IGF receptor mRNA
(% of GSV-treated)
125
100
75
50
25
0
0
30
60
90
Oligonucleotide concentration (nM)

Figure 33

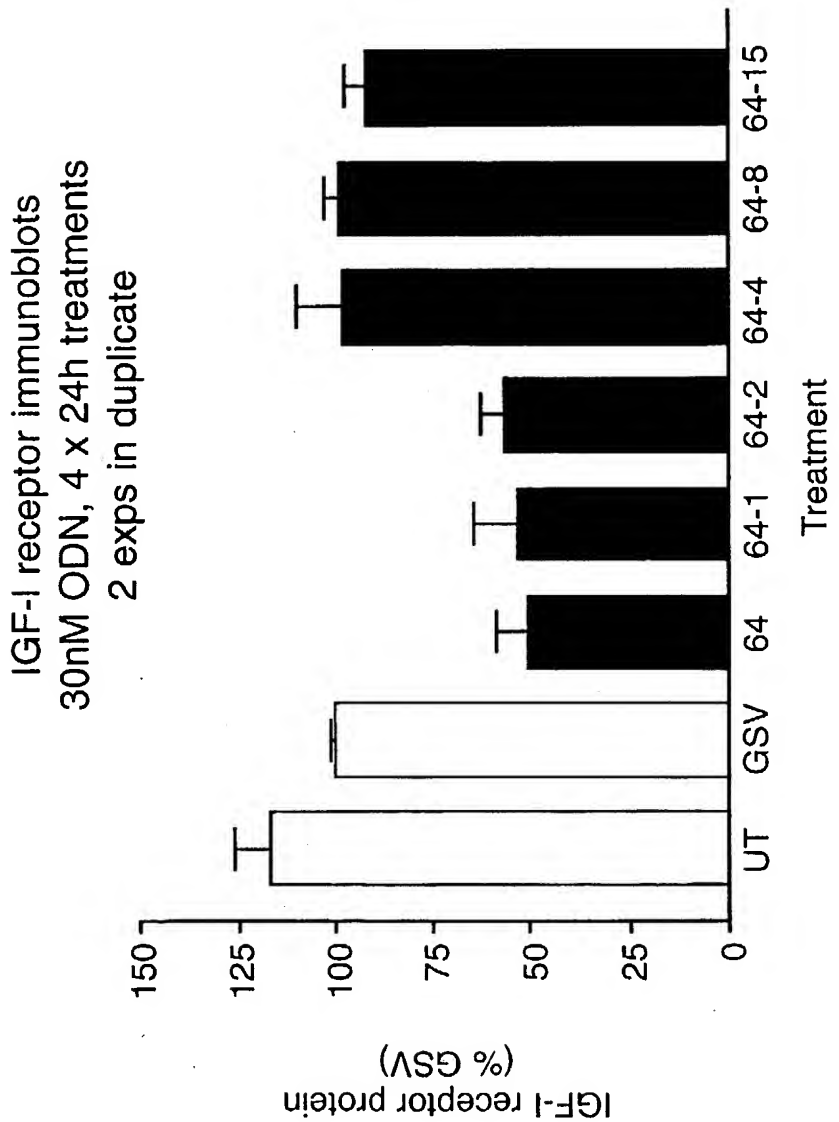

IGF-I receptor immunoblots
30nM ODN, 4 x 24h treatments
2 exps in duplicate
IGF-I receptor protein
(% GSV)
150
125
100
75
50
25
0
UT
GSV
64
64-1
64-2
64-4
64-8
64-15
Treatment

Figure 34

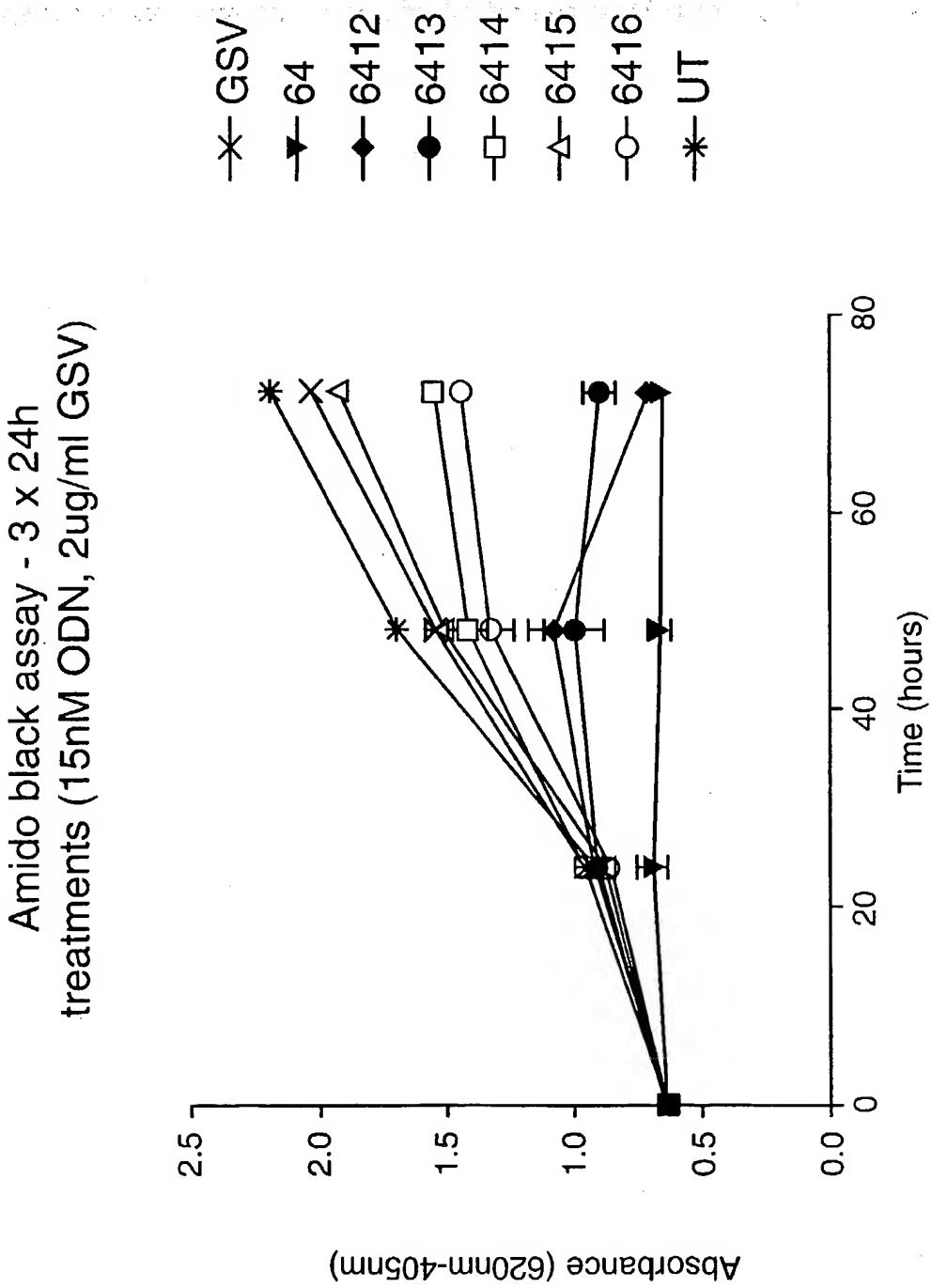

GSV
64
6412
6413
6414
6415
6416
UT
Amido black assay - 3 x 24h
treatments (15nM ODN, 2ug/ml GSV)
Absorbance (620nm-405nm)
2.5
2.0
1.5
1.0
0.5
0.0
0
20
40
60
80
Time (hours)

Figure 35

SEQUENCE LISTING

<110> MURDOCH CHILDREN'S RESEARCH INSTITUTE

<120> A METHOD FOR THE PROPHYLAXIS AND/OR TREATMENT OF MEDICAL DISORDERS

<130> 2288267/EJH

<140> INTERNATIONAL
<141> 2000-06-21

<150> 60/140345
<151> 1999-06-21

<160> 24

<170> PatentIn Ver. 2.1

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<213> synthetic construct

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gcgccgcccg ccgcggtggc cgcagtggcc ggaggcgccc gcatgccatg cgcggagctc 360
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aagcccctca agtcgggtat gaaggagctg gccgtgttcc gggagaaggt cactgagcag 720
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accatccggg gggaccccga gtgtcatctc ttctacaatg agcagcagga ggcttgcggg 1080
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gcccctctcc aaacaccggc agaaaacgga gagtgcttgg gtggtgggtg ctggaggatt 1200
ttccagttct gacacacgta tttatatttg gaaagagacc agcaccgagc tcggcacctc 1260
cccggcctct ctcttcccag ctgcagatgc cacacctgct ccttcttgct ttccccgggg 1320
gaggaagggg gttgtggtcg gggagctggg gtacaggttt ggggaggggg aagagaaatt 1380
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<211> 2474
<212> DNA
<213> synthetic construct

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gccgggctgc ggctgctgcc tgacgtgcgc actgagcgag ggccagccgt gcggcatcta 360
caccgagcgc tgtggctccg gccttcgctg ccagccgtcg cccgacgagg cgcgaccgct 420
gcaggcgctg ctggacggcc gcgggctctg cgtcaacgct agtgccgtca gccgcctgcg 480
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cagcgccggc agtgtggaga gcccgtccgt ctccagcacg caccgggtgt ctgatcccaa 600
gttccacccc ctccattcaa agataatcat catcaagaaa gggcatgcta aagacagcca 660
gcgctacaaa gttgactacg agtctcagag cacagatacc cagaacttct cctccgagtc 720
caagcgggag acagaatatg gtccctgccg tagagaaatg gaagacacac tgaatcacct 780
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attttataag aaaaagcagt gtcgcccttc caaaggcagg aagcggggct tctgctggtg 900
tgtggataag tatgggcagc ctctcccagg ctacaccacc aaggggaagg aggacgtgca 960
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actattggag aaaataaggt ggagtcctac ttgtttaaaa aatatgtatc taagaatgtt 1620
ctagggcact ctgggaacct ataaaggcag gtatttcggg ccctcctctt caggaatctt 1680
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aacattgtat acaacatagc cccaaatata gtaagatcta tactagataa tcctagatga 2160
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aagtttttac catt 2474

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<213> synthetic construct

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tggccgacga gtggagaaat ctgcgggcca ggcatcgaca tccgcaacga ctatcagcag 180
ctgaagcgcc tggagaactg cacggtgatc gagggctacc tccacatcct gctcatctcc 240
aaggccgagg actaccgcag ctaccgcttc cccaagctca cggtcattac cgagtacttg 300
ctgctgttcc gagtggctgg cctcgagagc ctcggagacc tcttccccaa cctcacggtc 360
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aatgctgacc tctgttacct ctccactgtg gactggtccc tgatcctgga tgcggtgtcc 540
aataactaca ttgtggggaa taagccccca aaggaatgtg gggacctgtg tccagggacc 600
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Cys Asp Lys His Gly Leu Tyr Asn Leu Lys Gln Cys Lys Met Ser Leu
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Asn Gly Gln Arg Gly Glu Cys Trp Cys Val Asn Pro Asn Thr Gly Lys
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Pro Gly Cys Gly Cys Cys Leu Thr Cys Ala Leu Ser Glu Gly Gln Pro
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Cys Gly Ile Tyr Thr Glu Arg Cys Gly Ser Gly Leu Arg Cys Gln Pro
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Ser Pro Asp Glu Ala Arg Pro Leu Gln Ala Leu Leu Asp Gly Arg Gly
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Leu Cys Val Asn Ala Ser Ala Val Ser Arg Leu Arg Ala Tyr Leu Leu
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Pro Ala Pro Pro Ala Pro Gly Asn Ala Ser Glu Ser Glu Glu Asp Arg
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Ser Ala Gly Ser Val Glu Ser Pro Ser Val Ser Ser Thr His Arg Val
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/AU 00/00693

A. CLASSIFICATION OF SUBJECT MATTER

Int Cl[7]: A61K 38/30; 17/06; 17/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Derwent WPAT IGF-1, IGFBP, Insulin Like Growth Factor/Binding Pair.

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	AU 77314/94,A (688793) (Celtrix Pharmaceuticals, Inc.) 30 March 1995.	1-9, 20-22, 25, 29-37
X	AU 28753/95,A (692278) (Royal Children's Hospital Research Foundation, Australia.) 25 January 1996.	1-13, 20-23, 29-36
X	Wraight, Christopher J. et al., Expression of insulin-like growth factor binding protein-3 (IGFBP-3) J. Invest. Dermatol. (1997), 108(4), 452-456.	1-13,20-23,29-36

[X] Further documents are listed in the continuation of Box C

[ ] See patent family annex

* Special categories of cited documents:

"A" Document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
24 August 2000	-4 OCT 2000
Name and mailing address of the ISA/AU	Authorized officer
AUSTRALIAN PATENT OFFICE PO BOX 200 WODEN ACT 2606 AUSTRALIA E-mail address: pct@ipaustralia.gov.au Facsimile No.: (02) 6285 3929	A. WILCOX Telephone No.: (02) 6283 2243

INTERNATIONAL SEARCH REPORT

International application No.
PCT/AU 00/00693

C (Continuation). **DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	Jeschke, Marc G.; Barrow, Robert E.; Hawkins, Hal K.; Chrysopoula, Mina S. T.; Perez-Polo, J. Regina; Herdon, David, N. Effect of Multiple gene transfers of insulin like growth factor I complementary DNA gene constructs in rats after thermal injury. Arch. Surg. (1999), 134(10), 1137-1141.	1-13,20-23,29-36
X	WO 96/01636 (Royal Childrens Hospital Research Foundation) 25 January 1996.	1-13
X	WO 96/33216 (Pharmacia AB) 24 October 1996.	1-13

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/AU 00/00693

Box 1 Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 1,2,5-8 have been partially searched.
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

The specification provides support for methods and compositions which inhibit IGF-1 mediated cell proliferation and/or inflammation. There is no basis in the specification for methods and compositions derived from an invention for the treatment of cell proliferation and/or inflammation mediated by factors other than IGF-1.

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a)

Box II Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:
The claims are directed to methods and compositions which inhibit IGF-1 mediated cell proliferation and/or inflammation. The broader claims include cell proliferation and/or inflammation mediated by keratinocyte growth factor (KGF), TGF-α, TNF-α, IL-1, IL-2, IL-6, IL-8 and/or basic fibroblast growth factor (bFGF). Claims 1,2 and 5-8 are considered to include multiple inventions.

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims

2. ☒ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.